

Client: Vitco, Inc.

#### WASTE CHARACTERIZATION ANALYSIS REPORT

Sample Description

A	ttn:	Nappanee Mr. Bret	IN 465!	50 mann	EIS Analys	is No.: 86% Pretre		ste
D:	ate Sa	mpled:	2-08-88		Composite.		Sweepings	
C	ollect	ed By:	Vitco			200		
D	ate Re	ceived:	2-17-88					
D	ate Fo	rwarded:	4-04-88					
P	urchas	e Order:	10341					
ana	lysis the ev	procedure aluation	s. The f procedure	ollowing s herein		ere utiliz	ed, as nee	saea,
•	"Test Metho	Methods ods" USEPA	for the E SW-846,	valuatio November	n of Solid Wa 1986, 3rd Ed	ste - Phys ition	ical/Chemi	cal
	"Meth	nods for C	hemical A	nalysis	of Water and	Wastes" EP	A 600/4-79	9-020
		e of India						
The wer	speci e the	ific clien following	t request	ed analy	sis for the s	amples des	cribed abo	ove
	EP To	oxicity -	Metals		tate of India			
	EP T	oxicity -	Organics		olatile Organ			
	Igni	tability			emi-Volatile			
	Corr	osivity		(	Base/Neutrals			
	Reac	tivity	•	<u>√</u> ₽	CB	Pesticid	les	
	Addi	tional						

Materials constituting this report packet include laboratory analysis bench sheets. These bench sheets are required by the State of Indiana as an integral part of the Waste Characterization Analysis Report. Certain sections of this report may not pertain to your samples but do constitute a part of the EIS Report Packet. All results are hand entered to eliminate data transfer errors.

Andris Rozite, Laboratory Director

RECEIVED APR 06 1988

Rev (12-09-87)

# QUALITY ASSURANCE DATA EP TOXICITY and/or LEACHING METHOD

Parameter	<pre>% Recovery USEPA EMSL</pre>	- Accuracy	% RSD ' Precision
F & L & L & L & L & L & L & L & L & L &	QC Sample	Matrix Spike	Analysis
Arsenic *	94.		
Barium *	108.9	-	
Cadmium *	101.	-	
Chromium *	106.	-	
Copper	101.6		
Iron			
Lead *	94.1	-	
Manganese			
Mercury *	99.2	-	
Nickel	103.1		
Selenium *	104.6		
Silver *	104.8		
Sodium			
Zinc	101.3		
Chlorides			
Cyanide, Total			
Fluoride	13 10 10 10 10 10 10 10 10 10 10 10 10 10		
PCB			
рН			
Phenols			
Sulfate			
Sulfide, Total	:		
TDS	10000000		
TOC			
TOH			

<sup>\*</sup> These metals are analyzed by the Method of Standard Additions

	EP Toxicity			a Leaching M te Types Onl	
Parameter	RCRA	A	В	. <u> </u>	D
Arsenic *	5.0	0.05	0.5	1.25	5.0
Barium *	100	1.	10.	25.	100.
Cadmium *	1.0	0.01	0.1	0.25	1.0
Chromium *	5.0	0.05	0.5	1.25	5.0
Lead *	5.0	0.05	0.5	1.25	5.0
Mercury *	0.2	0.002	0.02	0.05	0.2
Selenium *	1.0	0.01	0.1	0.25	1.0
Silver *	5.0	0.05	0.5	1.25	5.0
Chlorides	<b>NATION</b>	250.	2500.	6250.	**
Copper		0.25	2.5	6.25	**
Cyanide, Total	<del></del>	0.2	2.	5.	**
Fluoride		2.4	24.	60.	**
Iron	- -	1.5	15.	**	**
Manganese	-	0.05	0.5	**	**
Nickel	-	0.2	2.	5.	**
PCB		==	-		_
рН	-	6.0 - 9.0	5 - 10	4 - 11	**
Phenols	- ,	0.3	3.	7.5	**
Sodium		250.	2500.	6250.	**
Sulfate	east	250.	2500.	6250.	* *
Sulfide, Total	-	1.	5.	12.5	**
TDS	<b>e</b> ad	500.	5000.	12500.	**
TOC	<b>a</b> nd	444		_	MED
TOH	**	***			===
Zinc	ipus	2.5	25.	62.5	**

<sup>\*</sup> Limits shown are based on EP Toxicity Analysis Data

<sup>\*\*</sup> Testing is not required

# ANALYTICAL REPORT SHEET EP TOXICITY - METALS ANALYSIS

EIS Lab Number	628H								
Client Description	Composite 86% Pretreatment	Į.	•						
	14% Floor Sweepings								
* * * * * * * * * * * *	* * * * * *	* *	<b>*</b> *	* *	* *	* *	* *	* *	* * *
% Solids	100						· · . · . · . · . · . · . · . · . ·		
Weight Raw Sample (g)	100.1					5			
Filters Used	AP15								
	HAWP			·					
* * * * * * * * * * * * *	* * * * * *	* *	k *	* *	* *	* *	* *	* *	* * *
Initial Extract pH	9.8								
Final Extract pH (24 hr)	7.2								
Acid Added (24hr)(ml)	400.4								
Final Extract pH (28hr)				•					
Acid Added (4hr)(ml)				,					
Total Acid Added (ml)	400.4								
Total DI Water Added (ml)	1602								
Original Liquid Phase(ml)	0								
Final Extract Volume (ml)	2002.								* * *
* * * * * * * * * * * * * * * * * * * *	* * * * * *	* *	* *	* *	* *	* *	* *	* *	* * *
RCRA Metals		<u> </u>			ļ· ·		· ·		
Arsenic (PPM)	<0.01								
Barium (PPM)	<0.5			and Carbolia .					
Cadmium (PPM)	0.05							ļ	
Chromium (PPM)	0.06							ļ	
Lead (PPM)	<0.Z							<u> </u>	
Mercury (PPM)	<0.005								
Selenium (PPM)	<0.005		2						
Silver (PPM)	<0.05								

Note: Tables of pH adjustments with time have been extracted from the EIS sample work sheets and are reproduced in the section of this report containing laboratory bench sheets.

### ANALYTICAL REPORT SHEET

# ADDITIONAL TESTS PERFORMED ON EP TOXICITY EXTRACT

EIS Lab Number				8 /											*****						,
Client Descript	ion	86% ·	Preti Floor 2 - 8	ه الحراث الما الما كا الما إلى ا	entung	ś					-					]					
* * * * * * * *	* * *	*	* *	* *	* *	*	*	*	*	* *	*	*	*	*	*	*	*	*	*	*	* *
Weigh Raw Sampl	.e (g)															_					
Filters Used						-					-					_					
			<u>.                                    </u>			-	····				-	•				_					
						-					-					-	<del></del>				
DI Water Added	(ml)						******								·						
* * * * * * * *	· * - * *	*	* +	* *	* *	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	* *
PARAMETERS						_					_ _					_					
Copper	(ppm)		0.	04	<u>L</u>	_											ļ.——				
Iron	(ppm)					_					_										
Manganese	(ppm)																				
Nickel	(ppm)	:	12.	.7																	
Sodium	(ppm)																				
Zinc	(ppm)		1.	. 6																	
Chlorides	(ppm)										_										
Cyanide, Total	(ppm)									<u></u>											
Fluoride	(ppm)									,a_	_ _										
PCB	(ppm)			······································	·		<del></del>				_										,
Phenols	(ppm)										_						ļ				
Sulfate	(ppm)				***						_						-				
Sulfide, Total	(ppm)																				
TDS	(ppm)										_ _						-				
TOC	(ppm)					_					_ _										
тон	(ppm)	\									_ -						_				
ph after 24 ho	ours																				112

Note: The reverse side of this sheet lists reference methods utilized

### LEACHING METHOD ANALYTICAL REFERENCES

# LEACHATE FORMATION

The leaching procedure utilized, and the equipment employed, is described on the report sheet titled SUPPLEMENTARY DATA EXTRACTION PROCEDURE TOXICITY TEST. The only deviation from the description is that NO PH adjustments were made.

### ANALYTICAL METHODS

PARAMETER	SW-846 Method #	EPA 600/4-79-020 Method #
Chlorides		325.3
Copper	3010 / 7210	
Cyanide, Total	9010	
Fluoride		340.2
Iron		236.1
Manganese		243.1
Nickel	3010 / 7520	
PCB	8080	
рН		150.1
Phenols		420.2
Sodium		273.1
Sulfate		374.4
Sulfide, Total	9030	
TDS		160.2
TOC	9060	
ТОН	9020	
Zinc	3010 / 7950	·

#### HAZARDOUS CHARACTERISTICS ANALYTICAL REPORT SHEET

This report sheet presents results of Corrosivity, Ignitability and Reactivity analysis if so required by the client. The following definitions apply to this analysis. The method of analysis is listed on the reverse side of this form.

- Corrosivity A solid waste exhibits the characteristic of Corrosivity if it is aqueous and has a pH of less that or equal to 2 (<) or greater than or equal to 12.5 (>)
- 2. Ignitability A solid waste exhibits the characteristic of ignitability if it is a liquid, other than an aqueous solution, containing less than 24% alcohol by volume, and has a Flash Point less than 140 °F (60 °C)
- A waste is classified as reactive if it exhibits any of the properties listed in 40 CFR 261.23. The property determined by laboratory analysis is related to the presence of Cyanide and Sulfide. The current USEPA Action level is: Cyanide 250 mg HCN/kg waste; Sulfide 500 mg H2S/kg waste

EIS Lab Number	Client Sample Description		Ignitability (°F)(%C)	Reactivit As Rec Cyanide	y (mg/kg) eived Sulfide
628H	86% Pretreatment 14% Floor Sweepings 2-8-88	9.2	> /77	1.0	2.0
					- Manual
					1942 TAXABAR SAN

#### METHOD OF ANALYSIS

The following procedural summaries are provided for clarification. The specific steps involved can be found in the following reference commonly referred to as SW-846: "Test Methods for Evaluating Solid Waste, Physical/Chemical Methods", USEPA, SW-846, November 1986, 3rd Edition.

### CORROSIVITY (SW-846 Method 9040)

- . Liquid Waste (Aqueous) are analyzed directly for pH using a commercial pH meter.
- . Solid wastes are analyzed for pH by preparing a 10% W/V solution of the waste in deionized water.

# IGNITABILITY (SW-846 Method 1010 or 1020)

The method of analysis consists of heating a sample at a slow, constant rate with continual stirring in a closed cup tester. The Flash Point is defined as the lowest temperature at which a test flame applied to the sample ignites the vapors above the sample.

REACTIVITY (SW-846 Method 9010 - Cyanide and Method 9030 - Sulfide)

- Cyanide is determined by refluxing of the sample under acid conditions with subsequent trapping of HCN gas in a Sodium Hydroxide scrubber. The scrubber contents are then analyzed either by titration or auto analyzer.
- Sulfide is determined by sweeping Hydrogen Sulfide gas out of an acidified sample using a Nitrogen atmosphere. The HS gas is trapped in dual scrubbers containing Zinc Acetate and then analyzed by titration with Sodium Thiosulfate.

### METHOD OF STANDARD ADDITIONS

#### METALS ANALYSIS

It is also possible to analyze a sample by the method of additions and to calculate the results, rather than plotting an analytical curve. The equation to be used for the calculation is shown below. The following three criteria must be met in order for the calculation to be valid:

- 1. All solutions are made to the same final volume
- A plot of concentration vs. absorbance is linear over the entire concentration range covered
- All readings are either in absorbance or in some value directly proportional to absorbance ("concentration")

$$C_{S} = \frac{C_{a} \cdot R_{s} \cdot V}{V_{s}(R_{a} - R_{s})}$$

where

- $C_s$  = concentration of the element of interest in the original sample solution, in  $\mu g/ml$
- $C_a$  = concentration of the element of interest in the standard solution used to "spike" the diluted sample solution, in  $\mu g/ml$
- $V_S$  = volume of original sample solution used, in ml
- $V_a$  = Volume of standard solution used, in ml
- $R_{S}$  = reading obtained for diluted sample solution
- $R_a$  = reading obtained for "spiked" sample solution

# / THOD OF STANDARD ADDITIONS FOR METALS

:IS # 628H

1ETAL Cu DATE	2-24-88 Ca	libration St	d. 5 Absor	rbance .17	/ / 3
Sample (ml) 3.50	Water (ml) 1.50	Std. (ml) Ø.@Ø	mg/l added 0.00	Result Ca .03	0.04
3.50	1.00	. 50	.50	.52	. 0437
3.50	.50	1.00	1.00	1.01	. 0437
3.50	0.00	1.50	1.50	1.51	.0436
	0.00	at a turber	* * *****	2000	
1ETAL Zn DATE	2-24-88 Ca	dibration St	d. 1 Abso	rbarice . 108	
Sample (ml)	Water (ml)	Std. (ml)	mp/l added	Result Ca	1c (1.6)
4.00	6.00	Ø. ØØ	ଡ. ପଡ	.64	ଡ. ଉପ୍ରତ
4. ØØ	5.00	1.00	. 10	. 74	1.5875
	4.00	2.00	.20	. 83	1.6282
4.00			. 30	. 93 . 93	1.6144
4.00	3.00	3.00	* 12Ki	. 73	1. 0144
					•
1ETAL Cd DATE	2-24-88 Ca	alibration St	d. 2 Abso		
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	Result Ca	1c (0.05)
3.50	1.50	0.00	~ Q. QQ	<b>.</b> Ø4	<b>ଡ. ଉଉଉଡ</b> ି
3.50	1.00	.50	.20	. 26	.0519
3,50	.50	1.00	. 40	. 45	. 0557
	ଷ୍ଟ ଷଷ	1.50	.60	.67	. 2544
3,50	עני יציעי	1:00	: C.40	101	
1ETAL Ni DATE	: 2-24-88 Ca	alibration St	d. 2 Abso	rbance .047	PLATE 128
//	Dimbonia (m.1)	Std. (ml)	mm/l added	Result Ca	1c $(12.7)$
Sample (ml)		0.00	0.00	. 47	0.0000
3.50	1.50	.50	.20	. 67	. 6481
3.50	1.00				.6403 (N
3.50	. 50	1.00	. 40	.88	* C-60 ( )- I
3.50	ଡ. ଉପ	1.50	.60	1.10	. 6327 🕚
1ETAL Pb DATE	2-25-88 Ca	alibration S	td. 5 Abso	rbance .034	
<b></b>	11 m. da m / v n 1 \	Std. (ml)	mg/l added	Result Ca	11c (20.2)
	Water (ml)	Ø. ØØ	Ø. ØØ	.13	0.0000
3.50	1.50			.61	. 1935
3.50	1.00	.50	.50		.1848
3.50	. 50	1.00	1.00	1.14	. 1809
3.50	ଡ.ଡଡ	1.50	1.50	1.67	" TOO3

1ETAL Cr DATE	2-27-88 Ca	libration S	td. 5 Abso	rbance .13	7 PLATE 131
Sample (ml) 3.50 3.50 3.50 3.50	Water (ml) 1.50 1.00 .50 0.00	Std. (ml) 0.00 .50 1.00 1.50	0.00 .50	Result . 04 . 49 . 94 1. 42	Calc 0.0000 .0556 .0552 .0542
1ETAL Ag DATE	3-01-88 Ca	libration S	td. 2 Abso	rbance .i	PLATE 138
Sample (m1) 3.50 3.50 3.50 3.50	Water (ml) 1.50 1.00 .50 0.00	Std. (ml) Ø.ØØ .50 i.ØØ 1.50	mg/l added 0.00 .20 .40 .60	Result .02 .20 .39 .60	Calc (0.05) 0.0000 .0232 .0232 .0220
1ETAL BA DATE	3-02-88 Ca	libration E	Std. 20 Abs	orbance .1	47 PLATE 140
Sample (ml) 3.50 3.50 3.50 3.50	Water (ml) 1.50 1.00 .50 0.00	Std. (ml) 0.00 .50 1.00 1.50	mg/l added 0.00 2.00 4.00 6.00	Result .30 2.50 4.53 6.66	Calc <0.5 0.0000 .3896 .4053 .4043

# METHOD OF STANDARD ADDITIONS FOR METALS One Calibration Standard Used

<u>Se 3-9-8</u>	<u> </u>	28 H C	alibration S	td. <u>0.020</u>	(<0.005)
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	Conc. (mg/l	.02(,1) \$ -0.002 -4(1.8-0)
4	16	0	0	_1	-1(1.8-0)
	12	4-	0.004	0.0013	
	g i	8	0,008	0.0034	- 02(100) = <0.002 .
	4	. /2.	0.012	0.0064	4(3.4)
	· /				4(6.4-0)
As 3-10-99	= EIS #	62 <i>8 H</i> c	alibration S	Std. <u>0.0200</u>	- (20.01)
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	i  Conc. (mg/l	) 1/4/
4	16	0	0	0	140,005
1	/2	4	0.004	0.0036	4(3.6-0)
	3	8	0.009	0.0058	.02(1)8 . 20.005
	4	12	0.012	0.0107	4(5.5-0)
v.					.02(.1)/2 = <0.005 4(10.7-0)
	EIS #	С	alibration :	Std	
· ·					
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	d   Conc. (mg/l	<u>)                                    </u>
	l				
		!	.   <u></u>		
		1	.		
		l	.		<del></del>
		_		<b>a</b> t. 1	
	EIS #	C	Calibration	Sta.	
Sample (ml)	Water (ml)	1 Std. (ml)	mg/l adde	d   Conc. (mg/l	_)
Dumpio ()			_		
<del></del>			.		····
·.			_1		<del></del>
		.	_	<b> </b>	····
	EIS #	(	Calibration	Std.	
	Water (ml)	1 Std. (m1)	mg/l adde	ed   Conc. (mg/)	1)_
Sample (ml)	March (mr)	1 2 2 2			
	_				
	_	1			<del>,</del>
	_				

### Mercury Worksheet

Sulfuric Acid (H2SO4)	_5	11-19-87	Pot.	Dichroma	ate	(K2Cr207)		
Nitric Acid (HNO3)	2.5	12-9	Sod.	Cl.,-Hydi	rox.	Hydróchl	oride	
Pot. Permanganate (KMnO	4) 4.5	2-17-88		·(NaCl I	NH201	н·HCl)	4.3	1-20
Pot. Persulfate (K2S2O8	_							

					,	
a second	Bottle #	Sample #	Vol. or Wt.	Re haightan	sult	
1	/	BI		ne igut		
2	2	BI	_	0		
3	6	0.05 ug Hg		4		
4	7	o. I sig Hg		8		· · · · · · · · · · · · · · · · · · ·
5	9	0.2 mg Hg		17		
5		0.3 mg Hg		30		
7	14	o. 4ugHg		39		
3	16	533H	) 50-	2		
<del></del>	21	0	50ml +0. leng Hag	12		)
<b>)</b>	27		50ml +0.20g Hg	24	(20.005)	
1	29		1 50 -1+0. 3ug Hg	34		
2		543H ·	) 50 W	1		
3	32	0	50ml +0.1mg Hg	11	20,005	<u>ノ</u>
1	33		50-1+0.2 ceg Hg	21		
5	35		50ml + 0.3cmg Hay	35		<u>,</u>
5	44	5444	toom	2	<0.00	05
7		545H	200/	2	<0.00	200
	45	546H	200.	2	<0.0	05
3 :	48	547H	Loom	5	0.00	006
-	83	ius378-4	100/	15	0.00159	99.2% le
J	86	uctual = 0.0016	1			
		A				

95% C.I. = 0.0011-0.0021

COMMENTS

# Mercury Worksheet

Sulfuric Acid (II2SO4)	Pot. Dichromate (K2Cr207)
Nitric Acid (HNO3)	Sod. ClHydrox. Hydrochloride
Pot. Permanganate (KMnO4)	·(NaCl NH2OH·HCl)
Pot. Persulfate (K2S2O8)	Stannous Chloride (SnCl2)

•				
1 145	Bottle #	Sample #	Vol. or Wt.	Result
1	100	548H	food	Leight man <0,0005
2	10 5	5494	tooml	3 40.0005
3	105	549H DUP	toom	3.5 <0.000\$
4	113	574H	25~/	2
5	114	580 H	) 50ml	2
6	118	0	( 50.1 +0.1eg Hg	9 (20.005)
7	129		50ml + 0,2 ceq Hay	21
8	133		50-110,3 eg Ha	32
9	141	600 H	) soul	5
10	176	0	soul + oileighty	14
11	179		Dul + 0, Zeg Hg	25
12	182		Dult or Bug Hay	37
1 3	184	628H	501	
14	217	0	soul +orleighty	10 (20.005)
L5			Don't + 0,2 mg Hg	19
 L6	233		Dul 10.3 eng the	32
1,7	238	530#	(8)	5 .843-
L8	244	530H	(2) 5	18 .379
19	250 263	511 H	1 ml	1
19. 20	270	511 H	1 ml + 0. 3 mg Hg	4
20	J 4/V	1		

COMMENTS	

		and the first company of the Market and the second		 111.
	D-1-	2-19-88	Analyst	 MW
Analysis	Date	<u> </u>	I III CALL	

# P-XYLENE READINGS

Trial #	Observed Value C (F)	Barometric Pressure (mmHg)	Corrected Flash Point C (F)	Average Value Corrected Flash	Quality Control Limit
1	81°	731	82.7°	<del>-</del>	26.2 - 27.8 Centigrade
2	80°	732	81.7°	82.2	79.2 - 82.5 Fahrenheit

# SAMPLE ANALYSIS

SAMPLE ANALYSIS								
1 #	Sample Type L=Liquid S=Solid	Observed Value C F	Barometric Pressure (mm Hg)	Corrected Flash Point C F	Comments and/or Precision data			
573H	<u></u>	171°	735	172.5				
62814	S	7 175°	730	>176.8°				
		,						
		<del></del>	<del></del>		•			

# CALCULATION FOR CORRECTED FLASH POINT

- · Using oF and mm Hg

- · Using oC and mm Hg
- CORRECTED FLASH = (Observed Value) + [0.06 (760 Barometric Pressure)]

  CORRECTED FLASH = (Observed Value) + [0.033 (760 Barometric Pressure)]

•	Date 2/18-19/88 Analyst CV EIS# 628#
2.	Sample & Date Vitco, Inc
3.	Description of Sample: 86% Vietreatment weste 14% Floor sweepings
4.	Type of Analysis: Metals X Organics Other
5.	Wt. of raw sample used for first separation (100 gm. minimum) /02/
6.	Filters Used Weight (gm.) Filters Used Weight (gm.)
	HAWP 202.9,
	AP15 102-8
7	TOTAL WEIGHT + =
3.	Total weight of solid material and filters after filtration
).	- Less tare weight of filters used (line 7)
) .	= Actual weight of material for extraction $\frac{00.1}{}$
•	If the solid residue on the filters is less than 0.5% of the original
	weight of the waste, the residue can be discarded and treat the liquid phase as the extract.  Result from line 10 ( ) $\times$ 100 =
)	If the result from the formula in 11 is greater than 0.5% solids, you
<b>.</b>	must go. through extraction. The total amount of distilled water to be used
•	in transferring material to the extractor and which is to be used in the
	extraction is obtained by the following formula:
	Weight from line 10 ( $\frac{100}{100}$ ) x 16 = $\frac{1602}{1002}$
•	The total amount of 0.5N Acetic acid which can be used in pH adjustment
•	is obtained by the following formula:
	Weight from line 10 ( (00, $\sqrt{}$ ) x 4 = $\sqrt{400.4}$
	With residue and distilled water in extraction container (glass Must be
	used if organics are to be analysed - line 4 ) agitate and measure pH.
	If pH is greater than 5.0 decrease to 5.0 $\frac{+}{-}$ 0.2 by adding 0.5N acetic acid
	The first Angree of the acceptance of the Tark and the second of the sec

The pH of the solution should be adjusted at 15, 30 and 60 minute
intervals, moving to the next longer interval if the pH'does not have
to be adjusted more than 0.5 pH units to bring it to $5.0^{\pm}0.2$ . This
adjustment Must be carried on for AT LEAST 6 HOURS. This time will be
longer if the sample is still changing by more than 0.5 pH units at
60 minute or less time intervals. Adjust pH only when it rises above 5.2
Clock time Time Interval pH Adjusted pH Acid Added (ml.) 400.4
9:30 .25 9.8 4.8 140.4
10:13 7.9 5.8 260
· · · · · · · · · · · · · · · · · · ·
- Nor
Final pH 7.2 Total Acid Added 400.4
If at the end of the 24hr. extraction period the pH of the solution is
not below 5.2 and the maximum amt. of acid has not been added (line 13)
the pH should be adjusted to $5.0^{\pm}$ 0.2 and the extraction continued
for an additional FOUR hours, during which the pH will be adjusted
at one hour intervals.
Clock Time Time Interval pH Adjusted pH Acid Added (ml.)

19.	Final results if additional extraction time was dictated by
	Step 18 Final pH Total Acid Added (ml) 400.4
20.	
	should be added to the extractor in the amount determined by
	the following equation:
	$V = \{(20)(W)-16(W)\}-A$
	$V = [(20)(\underline{100.1}) - 16(\underline{100.1})] - (\underline{400.4}) = \underline{0}$
	V = ml distilled water to be added
	W = weight in grams of solid charged to extractor (line 10)
	A = ml of 0.5N acetic acid added during extraction (line 17 or
-,	19)
21.	This liquid in the extractor is now filtered once again and
	the liquid obtained is combined with the liquid which was
	obtained in the first filtration. This combined sample is now
	ready for analysis
22.	Volume (ml) of liquid from first filtration
	$m_{a}$ to 1 $m_{a}$ (m1) of DI water added to sample $\frac{160}{2}$
	Total Volume (ml) of acid added to sample (line 19) $\frac{400.4}{2002.4}$
	Total Volume (ml) 2002.4
23.	Comments:
24.	Solid Sample Preparation
	X 1. Completely sifted through a 9.5 mm standard sieve
	2. A surface area per gram of material equal to or
	greater than 3.1 square centimeters
	3. Been cut, crushed, or ground to the point where it
	may pass through a 9.5 mm standard sieve
	4. Been subjected to the Structural Integrity Procedure
	3.
25.	Paint filter liquids test (Method 9095) if performed.
	Free Liquid: YESNO

# PH METER STANDARDIZATION

ALYST	DATE	TIME	TEMP.	ACTUAL / READS		MODE	RANGE	QC
cu)	25	8:05	19°	9.0 / 7.0	4.01/4-0	N		
av .	2-16	8:00	18°	7.0/7.0	401/4-0	N		
as	2-18	6:15	180	7.0/7.0	4.4/4.0	N		
w	2-19	11:30	210	7.0/7.0	40/40	N		-
w	2-23-88	8:17	190	20/20	4.01/4-0	N		
au	2-24-88	8:07	190	7.0/7.8	4.0/4.0	N		
42	2-29-88	2.48	220	20/7.0	4.01/4.0	N		
<u>w</u>	3-1-88	3:05	270	7.0 / 7.0	4.01/4.0	N	,	
av	3.2-88	8:05	200	7.0/7.0	4.01/4.0	N		
<u> </u>	3-3-88	11:45	210	7.0/7.0	4.0/ 4.0	N		
cul	3.4-89	5:35	220	78/7.0	4.01/4.0	N		
W	3-7-88	'	200	7.0/7.0	4.01/4.0	N		
w	3-8-88	a; 50	210	7.0/ 7.0	4.0/ /4.0	N		
$\overline{\omega}$	3-9-86	8:05	21°	7.0/ 7.0	4.0/4.0	N		
$\frac{\omega}{\omega}$	3-10-86	8:45	200	7.0/ 7.0	4.0/ 4.0	N		
<del></del>								
······································								
	1		. 1		•			

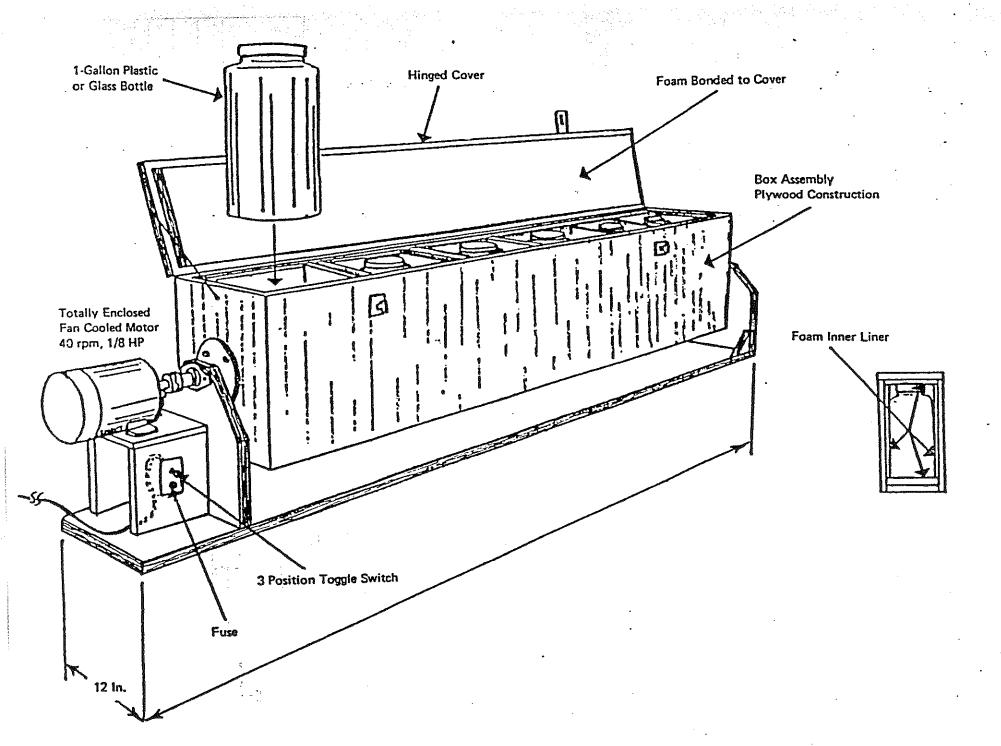
#### SUPPLEMENTARY DATA

#### EXTRACTION PROCEDURE TOXICITY TEST

- 1. The extractor used by EIS is shown in Figure 1 attached. The extractor meets the specifications of Method 1310, SW-846 "TEST METHODS FOR EVALUATING SOLID WASTE/PHYSICAL CHEMICAL METHODS" 2nd Edition, July 1982.
- 2. The filtration apparatus utilized by EIS is a positive pressure unit purchased from Millipore Corporation. The unit is Model No. YT30 142 HW and is rated at 100psi maximum inlet pressure. Nitrogen gas is used as the pressure source. All sample contact surfaces are Teflon coated.
- Filters utilized in the test procedure are purchased from Millipore Corporation.
   The specific filters utilized in this analysis are shown on the Analytical Report Sheet.
- 4. Test procedure used to form the leachate are exactly per specifications in Method 1310, SW-846. Measurement of pH is performed manually using a Fisher Accumet Model 420 pH Meter. Bench sheets showing pH adjustments with time are available for inspection at EIS. Samples are screened through a 3/8 inch sieve prior to filtration.
- 5. Metals analysis is performed using a Perkin-Elmer 560 Atomic Absorption Spectro-photometer for all metals except Mercury. Analysis of Mercury is performed using cold vapor technique on a dedicated Jarrell Ash Model No. 280 Atomsorb. Peripheral attachments for the 560 AA include Graphite Furnace with Auto Sampler, Hydride Generator and Auto Sampler for Flame Absorption.
- 6. All metals reported for EP Toxicity Analysis are analyzed by METHOD OF STANDARD ADDITIONS. This method is utilized after the extract has been digested per digestion procedures specified in SW-846.
- 7. The EIS data handling consists of calibrating the Atomic Absorption unit in terms of concentration values. Check standards are then analyzed along with samples. Output from the AA is hard copied by an Okidata 82A Microline Printer. Numerous client results are thus present on a single printer sheet. Subsequent submissions of bench type sheets to a client are made by manually extracting Method of Standard Additions data on to separate sheets.
- 8. The following SW-846 procedures were employed in this analysis.

	Digestion Procedure							
	Acid/	Acid/	Acid/	· Cold				
	Flame	Furnace	Hydride	Vapor	Other			
As		7060	7061					
Ba	3010	3020						
Cd	3010	3020						
Cr	3010	3020						
Pb	3010	3020						
Hg		!		7470				
Se		7740	7741					
Ag	7760	7761						

Analysis Procedure							
		•	Co1d				
Flame	Furnace	Hydride	Vapor				
	7060	7061					
7080	7081						
7130	7131						
7190	7191						
7420	7421						
		(	7470				
	7740	7741					
7760	7761						



# VITCO. INC.

#### P. O. BOX 407, NAPPANEE, INDIANA 46550

PHONE: 219-773-3181

September 21, 1987

Julie Tepe Prairie View Landfill 15505 Shively Road Wyatt, Indiana 46595

Dear Julie,

Please have the composite sample of porcelain enamel overspray and pretreatment waste collected 9-18-87 analyzed as soon as possible so we may begin disposal at your site. We have already been granted approval for disposal by the State of Indiana.

I have also enclosed a copy of the letter I sent to Sandy Jordan, IDEM, that explains the high and low levels of cadmium in the analysis of our composite samples in the past. The letter also explains how I intend to keep these levels of metals at a minimum in the waste destined for disposal at Prairie View.

Again, I hope this sample can be analyzed as soon as possible. Your cooperation thus far has been greatly appreciated! Please feel free to call me if you have any questions (219/773-3181).

Sincerely,

VITCO, INC.

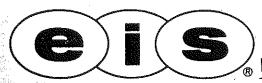
Butt L. Mordmann

Brett L. Nordmann Ceramic Engineer

BLN/vlc

cc : Len Micinski

enclosure



Vitco

Client:

# EIS ENVIRONMENTAL ENGINEERS, INC.

1701 North Ironwood Drive • South Bend, Indiana 46635 • 219/277-5715

Sample Description

#### WASTE CLASSIFICATION ANALYSIS REPORT

ATTN: Date Sampled: Date Received: 7-21-87 Date Forwarded: 7-28-87 Purchase Order: 84900	EIS Analysis No.: 2311G-2312G  2311G - Composite of Pretreatment Waste #1  2312G - Composite of Pretreatment Waste #2
This report presents results of analysis procedures. The followin the evaluation procedures he	waste classification through laboratory wing references were utilized, as needed, erein.
<ul> <li>"Test Methods for the Evaluation Methods" USEPA SW-846, July</li> </ul>	uluation of Solid Waste - Physical/Chemical uly 1982, 2nd Edition
• "Methods for Chemical Ana • State of Indiana "Leachir	alysis of Water and Wastes" EPA 600/4-79-020 ag Method"
The specific client requested a were the following.	analysis for the samples described above
EP Toxicity - Metals X	State of Indiana Leaching Method
EP Toxicity - Organics	Volatile Organic Compounds
Ignitability	Semi-volatile Organic Compounds
Corrosivity	(Base/Neutrals Acid Fraction)
Reactivity	PCB Pesticides
Additional	<u> </u>
bench sheets. These bench sheet	port packet include laboratory analysis are required by the State of Indiana te Classification Analysis Report. Certain

sections of this report may not pertain to your samples but do constitute a part of the EIS Report Packet. All results are hand entered to eliminate

Andris Rozite, Laboratory Director

data transfer errors.

# QUALITY ASSURANCE DATA EP TOXICITY and/or LEACHING METHOD

		- Accuracy	% RSD
	6 1.000	- Accuracy	Precision
Parameter	USEPA EMSL QC Sample	Matrix Spike	Analysis
Arsenic *		-	
Barium *		-	0 (23126)
Cadmium *	103.4	-	0 (23/26)
Chromium *		-	
Copper			
Iron			
Lead *			
Manganese			
Mercury *		-	
Nickel			
Selenium *			
Silver *	:		
Sodium			
Zinc	·		
Chlorides			
Cyanide, Total			
A.Fluoride	·		
PCB			
pH .			
Phenols			
Sulfate			
Sulfide, Total			
TDS			
TOC			
TOH		the Method of Sta	

\* These metals are analyzed by the Method of Standard Additions RECEIVEU JUL 30 1987

Sec. 1995.	TD Movielt	Stat	e of India	na Leaching N	Method
Parameter	EP Toxicity RCRA	A	Foundry Wa B	ste Types Onl	D
Arsenic *	5,.0	0.05	0.5	1.25	5.0
Barium *	100	1.	10.	25.	100.
Cadmium *	1.0	0.01	0.1	0.25	1.0
Chromium *	5.0	0.05	0.5	1.25	5.0
Lead *	5.0	0.05	0.5	1.25	5.0
Mercury *	0.2	0.002	0.02	0.05	0.2
Selenium *	1.0	0.01	0.1	0.25	1.0
Silver *	5.0	0.05	0.5	1.25	5.0
Chlorides	<del>(mp</del>	250.	2500.	6250.	**
Copper	NO.	0.25	2.5	6.25	**
Cyanide, Total	-	0.2	2.	5.	**
Fluoride	<b>424</b>	2.4	24.	60.	**
Iron	<del></del>	1.5	15.	**	**
Manganese	_	0.05	0.5	* *	**
Nickel		0.2	2.	5.	**
PCB	400	•	•••		-
PH	-	6.0 - 9.0	5 - 10	4 - 11	* *
Phenols	- (	0.3	3.	7.5	* *
Sodium	<del></del>	250.	2500.	6250.	* *
Sulfate	_	250.	2500.	6250.	* *
Sulfide, Total		1.	5.	12.5	* *
TDS	6997	500.	5000.	12500.	**
TOC	<b></b>	<del></del>	_	<del></del>	920
тон	-		-	ana.	_
Zinc	-	2.5	25.	62.5	**

<sup>\*</sup> Limits shown are based on EP Toxicity Analysis Data

<sup>\*\*</sup> Testing is not required

# ANALYTICAL REPORT SHEET EP TOXICITY - METALS ANALYSIS

IIS Lab Number	23116	23126		
Client Description	Composite of Pretreatment	Composite of Pretreatment Waste #2	'	
	waste #1	* * * * * *	* * * * *	* * * * *
* * * * * * * * * * * * * * * * * * *	NO FREE LIQUID	100		
Weight Raw Sample (g)	100.3	102.4		
Filters Used	AP15	AP15		
FIICEIS OSCA	HAWP	HAWP		
i ko	* * * * * *	* * * * *	* * * * * *	* * * *
* * * * * * * * * * * * * * * * * * *	10.7	10.6		
Final Extract pH (24 hr)	8.8	8.9		
Acid Added (24hr)(ml)	401.2	409.6		
Final Extract pH (28hr)	_			
Acid Added (4hr)(ml)				
Total Acid Added (ml)	401.2	409.6		
Total DI Water Added (ml)	1605.	1638.		
Original Liquid Phase(ml)	0	0		
Final Extract Volume (ml)		2047.6	* * * * *	* * * * *
* * * * * * * * * * * * * * * * * * *				
Arsenic (PPM)				
Barium (PPM)				
Cadmium (PPM)	40,05	20.05		
Chromium (PPM)				
Lead (PPM)				
Mercury (PPM)				
Selenium (PPM)				
Silver (PPM)			· * * * * *	* * * * *

Note: Tables of pH adjustments with time have been extracted from the EIS sample work sheets and are reproduced in the section of this report containing laboratory bench sheets.

Put your address in the RETURN TO" space on the revers card from being returned to you. The return receipt fee will delivered to and the date of delivery. For additional service postmaster for fees and check box(es) for additional service.  1. Show to whom delivered, date, and addressee's addressed to:  Sanly Jacker Management Brank bold + Hamilton Wester Management Brank 105 S. Mirichan Street Indianapolis, IN 46225  July  5. Signature – Addressee  X  Via 6. Signature – Agent X  7. Date of Delivery  AUG 0 3 1987	e-sido. Failure to do this will proprovide you the name of the profollowing services are available	NO INSUMAN NOT HAVE Sandy Jord Marchine Washington Washington Washington Washington Washington The Sandian Andrews Sandian And	Managements 46225 73 .75
Dep PS Form 3811, Feb. 1986 Sol 105 South Meridian Street Indianapolis, IN 46225	DOMESTIC RETUR		
		ဦးသမ္မွာ သွေးချီသေ	

Dear Sandy,

I have enclosed a copy of the laboratory report of two samples of pretreatment waste which you requested an analysis for cadmium under EP toxicity. The results show the levels of cadmium in both samples to be well below the RCRA limitation.

Please complete the approval of our special waste application at your earliest convenience. Once completed, please send all relevent information, permits, etc., to Robert Koehl, President, VITCO, Inc.

Your prompt attention will be greatly appreciated.

Sincerely,

VITCO, Inc.

Beet L. Nordmann

Brett L. Nordmann Ceramic Engineer

BLN/nkn

cc: Robert Koehl, VITCO, Inc.

enclosure

	Date 3-5/3-6-87 Analyst CB EIS# 653
2.	Sample & Date Vico Inc. 2-12-87
3.	Description of Sample: Sample C. Floor Sweepings (P.E. Overspran)
4.	Type of Analysis: Metals Organics Other
5.	Wt. of raw sample used for first separation (100 gm. minimum) 101.3
6.	Filters Used Weight (gm.) Filters Used Weight (gm.)
•	HAWP 204,0
	AP15
	101.3
	DRY
7.	TOTAL WEIGHT + =
8.	Total weight of solid material and filters after filtration
9.	- Less tare weight of filters used (line 7)
LO.	= Actual weight of material for extraction
1.	If the solid residue on the filters is less than 0.5% of the original
	weight of the waste, the residue can be discarded and treat the liquid
•	phase as the extract. Result from line 10 ( ) $\times$ 100 =
L2.	If the result from the formula in 11 is greater than 0.5% solids, you
	must go. through extraction. The total amount of distilled water to be used
	in transferring material to the extractor and which is to be used in the
	extraction is obtained by the following formula:
	Weight from line 10 (101.3 ) x 16 = $1621$
13.	The total amount of 0.5N Acetic acid which can be used in pH adjustment
	is obtained by the following formula:
	Weight from line 10 ( $101.3$ ) x 4 = $405.2$
1.4.	With residue and distilled water in extraction container (glass Must be
<i>*</i>	used if organics are to be analysed - line 4 ) agitate and measure pH.
	If pH is greater than 5.0 decrease to 5.0 $\frac{+}{2}$ 0.2 by adding 0.5N acetic acid

l.

The pH of the solution should be adjusted at 15, 30 and 60 minute intervals, moving to the next longer interval if the pH does not have to be adjusted more than 0.5 pH units to bring it to  $5.0^{\pm}0.2$ . This adjustment Must be carried on for <u>AT LEAST 6 HOURS</u>. This time will be longer if the sample is still changing by more than 0.5 pH units at 60 minute or less time intervals. Adjust pH only when it rises above 5.2

	OO MINGE				_		מ מיעני
16.	Clock time	Time Interval	рН	Adjusted pH	Acid Added	(m1.)	405,2
	8127	0.25	8.8	5,1	<u>35. / </u>		370.1
	91/09	0.25	6.1	5.0	-25.0	60.1	345.1
	0124	0.00	5	50	13.1	73.2	3320
	9:37	10	1 th	5.0	14.0	87.2	318.0
	10:15	10	5.4	5.0	25.0	112.2	293.0
	11:25	1.0	<u>5,3</u>		<u> </u>	1100.0~	<b>0</b> - 1- <b>0</b>
	12:30	1.0	6.1				
	1:34	/. 💍	5,1				
	2.38	1.0	5,2			<del>_</del>	
	3145	18.0	5.2				
	_0.70						
٠		0+P A+	9.46-	Service Control of the Control of th		<u> </u>	
17.		Final pH	<u>کک</u>	Total Acid Add	$\frac{112.}{}$	<u> </u>	

18. If at the end of the 24hr. extraction period the pH of the solution is not below 5.2 and the maximum amt. of acid has not been added (line 13) the pH should be adjusted to 5.0 to 0.2 and the extraction continued for an additional FOUR hours, during which the pH will be adjusted at one hour intervals.

it one nour	THEOTYGET			- 4 9 3	
Clock Time	Time Interval	рН Л	djusted pH	Acid Added (ml.)	293.0
JOCK TIME			C 6	217 1432	262.7
:9's~s~	10	<u>ح</u> ے <u>ح</u>	<u> </u>	5/11	00000
		~ ~			
10158	1.0	5.D -	<u> </u>		
10100	/ >	5 No			
12.05	1.0	0.00			
- 0/		MAC			
' / ) X	(, /)	$50^{\circ}$			

- Final results if additional extraction time was dictated by 19. Total Acid added (ml) \_\_\_/43.2\_ Final pH 5,/ At the end of the 24 hour extraction period, distilled water 20. should be added to the extractor in the amount determined by the following equation: V = (20)(W) - 16(W) - A $V = [(20)(\underline{101.3}) - 16(\underline{161.3})] - (\underline{143.2}) = \underline{262}$ V = ml distilled water to be added W = weight in grams of solid charged to extractor (line 10) A = ml of 0.5N acetic acid added during extraction (line 17 or 19) This liquid in the extractor is now filtered once again and the 21. liquid obtained is combined with the liquid which was obtained in the first filtration. This combined sample is now ready for analysis. Volume (ml) of liquid from first filtration 22. Total Volume (ml) of DI water added to sample Total Volume (ml) of acid added to sample (Line 19) COMMENTS: 23.
- 24. Solid Sample Preparation
  - 1. Completely sifted through a 9.5 mm standard sieve
  - 2. A surface area per gram of material equal to or greater than 3.1 square centimeters
  - 3. Been cut, crushed, or ground to the point where it may pass through a 9.5 mm standard sieve
  - 4. Been subjected to the Structural Integrity Procedure 3.

	Date 3-5/3-6-87 Analyst B EIS# 654
	Sample & Date Vitto Inc. 2-12-87
•	Description of Sample: Sample D Solid Paint Wasie (Point over Sp.
	Type of Analysis: Metals Organics Other
	Wt. of raw sample used for first separation (100 gm. minimum) 101.5
	Filters Used Weight (gm.) Filters Used Weight (gm.)
	HAWP 201.4
	AP15
***	DR4
	TOTAL WEIGHT +
	Total weight of solid material and filters after filtration
) .	- Less tare weight of filters used (line 7)
} •	= Actual weight of material for extraction
	If the solid residue on the filters is less than 0.5% of the original
	weight of the waste, the residue can be discarded and treat the liquid
÷	Phase as the extract. Result from line 10 ( $\times$ X 100 = $\times$ Weight from line 5 ( $\times$ X 100 = $\times$
! <b>•</b>	If the result from the formula in 11 is greater than 0.5% solids, you
	must go. through extraction. The total amount of distilled water to be used
	in transferring material to the extractor and which is to be used in the
	extraction is obtained by the following formula:
٠.	Weight from line 10 ( $101.5$ ) x 16 = $1624^{\circ}$
•	The total amount of 0.5N Acetic acid which can be used in pH adjustment
	is obtained by the following formula:
	Weight from line 10 ( $101.5$ ) x 4 = $406.0$
•	With residue and distilled water in extraction container (glass Must be
	used if organics are to be analysed - line 4 ) agitate and measure pH.
	If pH is greater than 5.0 decrease to 5.0 $\frac{+}{-}$ 0.2 by adding 0.5N acetic acid.

6.

The pH of the solution should be adjusted at 15, 30 and 60 minute intervals, moving to the next longer interval if the pH does not have to be adjusted more than 0.5 pH units to bring it to  $5.0^{\pm}0.2$ . This adjustment Must be carried on for AT LEAST 6 HOURS. This time will be longer if the sample is still changing by more than 0.5 pH units at 60 minute or less time intervals. Adjust pH only when it rises above 5.2

	60 minute or	Tess time inco		_			
	Clock time	Time Interval	рН_	Adjusted pH	Acid Added	1 (ml.)	406.0
		1 20	6.1	4.9_	2.0		404.0
	8:37	0.20	69	< N	2.2	4.2	401.8
a' .	9:09	0.25	011	1/ 9	2.8	7.0	399.6
d i	9:37	0.50	<u>ک,ک</u>	4.1			396.0
:	10:15	6.3	5.4	4.9	3.0	10.0	391.0
		1,0	5.5	- 5,0_	. <u>5.0</u>	15.D	
	11125		<u>5,3</u>	4.95	5.0	20.0	386.0
	12:30	100		1/ 0	6.0	26.0	380.0
	1:34		<u>5,5</u>	3 _ 4.9	0.0		
	2:38	(0	5.2				370.C
	3:45	18.0	53	4.9	10.0	36.0	
	3.70			•			
			<u> </u>	<del></del>		•	
	<del></del>					<del>_</del>	
			3.1.4		· ~ /	۶ کا	

17. Final pH 5.9 Total Acid Added 36.0

18. If at the end of the 24hr. extraction period the pH of the solution is not below 5.2 and the maximum amt. of acid has not been added (line 13) the pH should be adjusted to  $5.0^{\pm}$  0.2 and the extraction continued for an additional <u>FOUR</u> hours, during which the pH will be adjusted at one hour intervals.

at one hour	intervais.	•	• • • • • • • • • • • • • • • • • • • •	Acid Added (ml.)	3703
Clock Time	Time Interval	pH C G	Adjusted pH	20.D 56.D	350
9:55	1.0	<u>5.7</u> 5.1			340.
12:05	1.0	5.25	5.0	10.0 66.0	340.
1:08		5.2			

7.

19.	Final results if additional extraction time was dictated by Step 18 Final pH 5.2 Total Acid added (ml) 66.0
	At the end of the 24 hour extraction period, distilled water

should be added to the extractor in the amount determined by the following equation:

$$V = [(20) (W) - 16 (W)] - A$$

$$V = [(20) (101.5) - 16 (101.5)] - (166.0) = 340$$

V = ml distilled water to be added

W = weight in grams of solid charged to extractor (line 10)

A = ml of 0.5N acetic acid added during extraction (line 17 or 19)

- 21. This liquid in the extractor is now filtered once again and the liquid obtained is combined with the liquid which was obtained in the first filtration. This combined sample is now ready for analysis.
- 22. Volume (ml) of liquid from first filtration

  Total Volume (ml) of DI water added to sample 1964

  Total Volume (ml) of acid added to sample (Line 19)

23.	COMMENTS:

24. Solid Sample Preparation

<b>✓</b>	l.	Completely sifted through a 9.5 mm standard sieve
	2.	A surface area per gram of material equal to or greater

than 3.1 square centimeters

Reen cut, crushed, or ground to the point where it may

3. Been cut, crushed, or ground to the point where it may pass through a 9.5 mm standard sieve

4. Been subjected to the Structural Integrity Procedure 3.

# PH METER STANDARDIZATION

٠.			:			1		
VALYST	DATE	TIME	TEMP.	ACTUAL / READS	ACTUAL / READS	MODE	RANGE	QC
CB_	2-27-87	8:15	20°	7.0/7.0	4.014.0	Μ		
	3-3-87	ł	1	7.0/7.0	4.014.0	M		_
B	3-4-87	10:12	210	7.0/ 7.0	4.01/4.0	N		
<u>CB</u>	3-5-87	7:50	210	7.0/7.0	4.01 / 4.0	N		-
<u>us</u>	3-6-87	8:29	20°	7.0/7.0	4.01/4.0	<b>/</b>		
CB	3-11-87	7:56	210	7.017.0	4.01/4.0	N		
_CB_	3-12-87	8:1/	20°	7.0/7.0	4.01/4.0	N		
<u>CB</u>	3-16-87	7:55	200	7.0/7.0	4.01/4.0	Λ/	'	
CB	3-17-87	9;37	200	7.0/7.0	4.01/4.0	N		
CB	3-18-87	8:02	220	7.0/7.0	4.01/4.6	N		
CB	3-19-87	8:21	210	7.0 /7.0	4.01/4.0	N		,
CB_	3-20-87	8:08	190	7.0/7.0	4.01/4.0	N		1
СВ	3-23-87	8:58	20°	70/7.0	4.01/4.0	N		
CB	3-24-87	8:22	210	7.0/7.0	4.01/4.0	N		
CB	3-30-87	7:58	190	7.017.0	4.01/4.0	N		
•								
<u> </u>								

#### SUPPLEMENTARY DATA

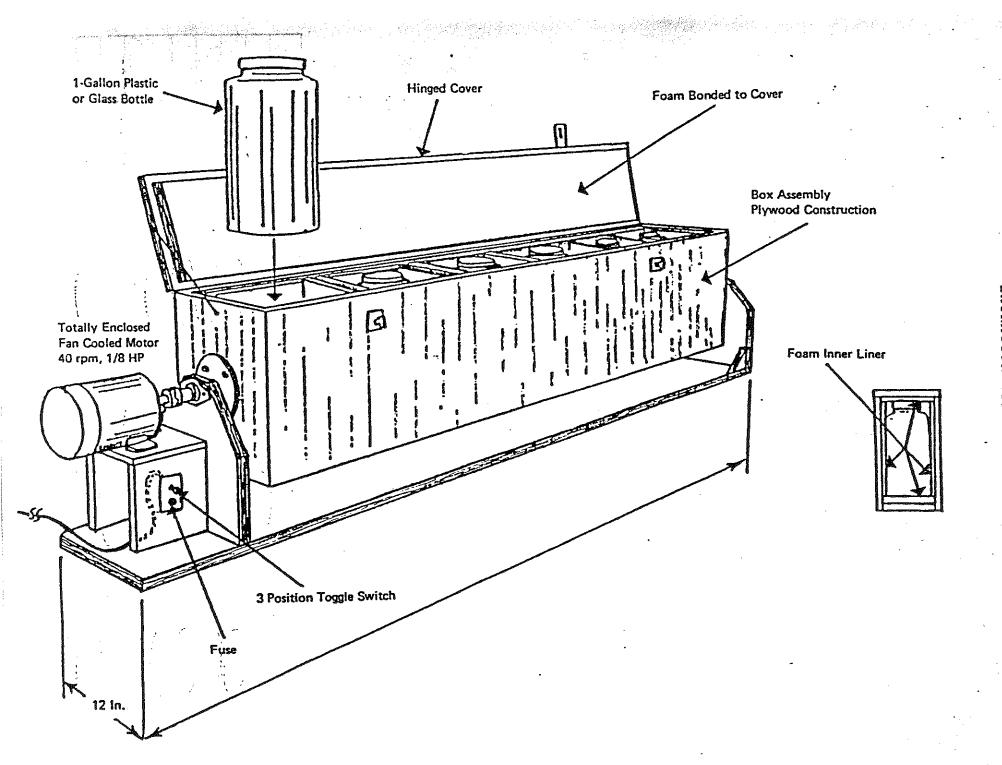
### EXTRACTION PROCEDURE TOXICITY TEST

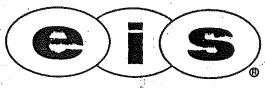
- 1. The extractor used by EIS is shown in Figure 1 attached. The extractor meets the specifications of Method 1310, SW-846 "TEST METHODS FOR EVALUATING SOLID WASTE/PHYSICAL CHEMICAL METHODS" 2nd Edition, July 1982.
- 2. The filtration apparatus utilized by EIS is a positive pressure unit purchased from Millipore Corporation. The unit is Model No. YT30 142 HW and is rated at 100psi maximum inlet pressure. Nitrogen gas is used as the pressure source.
  All sample contact surfaces are Teflon coated.
- 3. Filters utilized in the test procedure are purchased from Millipore Corporation.

  The specific filters utilized in this analysis are shown on the Analytical

  Report Sheet.
- 4. Test procedure used to form the leachate are exactly per specifications in Method 1310, SW-846. Measurement of pH is performed manually using a Fisher Accumet Model 420 pH Meter. Bench sheets showing pH adjustments with time are available for inspection at EIS. Samples are screened through a 3/8 inch sieve prior to filtration.
- 5. Metals analysis is performed using a Perkin-Elmer 560 Atomic Absorption Spectro-photometer for all metals except Mercury. Analysis of Mercury is performed using cold vapor technique on a dedicated Jarrell Ash Model No. 280 Atomsorb. Peripheral attachments for the 560 AA include Graphite Furnace with Auto Sampler, Hydride Generator and Auto Sampler for Flame Absorption.
- 6. All metals reported for EP Toxicity Analysis are analyzed by METHOD OF STANDARD ADDITIONS. This method is utilized after the extract has been digested per digestion procedures specified in SW-846.
- 7. The EIS data handling consists of calibrating the Atomic Absorption unit in terms of concentration values. Check standards are then analyzed along with samples. Output from the AA is hard copied by an Okidata 82A Microline Printer. Numerous client results are thus present on a single printer sheet. Subsequent submissions of bench type sheets to a client are made by manually extracting Method of Standard Additions data on to separate sheets.
- 8. The following SW-846 procedures were employed in this analysis.

Digestion Procedure							Analysis Procedure				
	Acid/ Flame	Acid/ Furnace	Acid/ Hydride	Cold Vapor	Other		Flame	Furnace	Hydride	Cold Vapor	
As		7060	7061			֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֡֓֓֓֓֓֡֓֓֡֓֡		7060	(7061)		
Ba	3010	3020				(	7080	7081			
Cd	3010	3020					7130	7131			
	3010	3020				1	7190	7191			
Cr		3020				1	7420	7421			
Pb	3010	3020	<del> </del>	7470	5	1			(	7470	
Hg		77/0	7741		1	1		7740	7741	<u>}</u>	
Se		7740	1/41	1		1	7760	7761			
Ag	(7760)	7761			<u> </u>	_1					





## EIS ENVIRONMENTAL ENGINEERS, INC.

1701 North fronwood Drive • South Bend, Indiana 46635 • 219/277-5715

#### WASTE CLASSIFICATION ANALYSIS REPORT

Sample Description  EIS Analysis No.: 651G - 654G  651G - Sample A; Composite of Pretreatment Waste 652G - Sample B; Paper Filter from Production Line 653G - Sample C; Floor Sweepings 654G - Sample D; Solid Paint Waste with Filters & Paint Overspray
f waste classification through laboratory owing references were utilized, as needed, erein.  aluation of Solid Waste - Physical/Chemica uly 1982, 2nd Edition  alysis of Water and Wastes" EPA 600/4-79-0
ng Method"  analysis for the samples described above  _ State of Indiana Leaching Method  Volatile Organic Compounds  Semi-volatile Organic Compounds  (Base/Neutrals Acid Fraction)
(

Materials constituting this report packet include laboratory analysis bench sheets. These bench sheets are required by the State of Indiana as an integral part of the Waste Classification Analysis Report. Certain sections of this report may not pertain to your samples but do constitute a part of the EIS Report Packet. All results are hand entered to eliminate data transfer errors.

Andris Rozite, Laboratory Director

# QUALITY ASSURANCE DATA EP TOXICITY and/or LEACHING METHOD

	% Recovery	- Accuracy	% RSD
Parameter	USEPA EMSL QC Sample	Matrix Spike	Precision Analysis
	QC Sample	HACLIX DPIKE	711141 7 5 1 5
Arsenic *	95.7	-	
Barium *	108.8	0.00	0 (6526)
Cadmium *	108.	•	2.7 (6526)
Chromium *	99.8	-	0 (6526)
Copper			
Iron	·		
Lead *	94.	-	0 (6524)
Manganese			
Mercury *	102.2 , 120.8	-	
Nickel			
Selenium *	1/3.8		
Silver *	104.8	-	
Sodium			
Zinc			
Chlorides			
Cyanide, Total			
Fluoride			
PCB			
рН			
Phenols			
Sulfate			
Sulfide, Total			
TDS			
TOC			
ТОН			

<sup>\*</sup> These metals are analyzed by the Method of Standard Additions

	EP Toxicity	State of Indiana Leaching Method Foundry Waste Types Only				
Parameter	RCRA	A	В	C	D	
Arsenic *	5.0	0.05	0.5	1.25	5.0	
Barium *	100	1.	10.	25.	100.	
Cadmium *	1.0	0.01	0.1	0.25	1.0	
Chromium *	5.0	0.05	0.5	1.25	5.0	
Lead *	5.0	0.05	0.5	1.25	5.0	
Mercury *	0.2	0.002	0.02	0.05	0.2	
Selenium *	1.0	0.01	0.1	0.25	1.0	
Silver *	5.0	0.05	0.5	1.25	5.0	
Chlorides	889	250.	2500.	6250.	**	
Copper	-	0.25	2.5	6.25	* *	
Cyanide, Total	u	0.2	2.	5.	**	
Fluoride	•••	1.4	14.	35.	**	
Iron	<b>S</b> ALES	1.5	15.	**	**	
Manganese		0.05	0.5	**	**	
Nickel		0.2	2.	5.	**	
PCB	_	-	==	-	<b>2002</b>	
рH	-	6.0 - 9.0	5 - 10	4 - 11	**	
Phenols	_	0.3	3.	7.5	**	
Sodium		250.	2500.	6250.	**	
Sulfate	<b>-</b>	250.	2500.	6250.	**	
Sulfide, Total	_	_	5.	12.5	**	
TDS	_	500.	5000.	12500.	* *	
TOC	-	_		, end	concr.	
TOH	· ·	_			_	
Zinc	-	2.5	25.	62.5	**	

<sup>\*</sup> Limits shown are based on EP Toxicity Analysis Data

<sup>\*\*</sup> Testing is not required

## ANALYTICAL REPORT SHEET EP TOXICITY - METALS ANALYSIS

EIS Lab Number	6516	6526	6 <i>53</i> 6	6546
Client Description	Sample A Composite of Pretreatment Waste	Sample B Paper Filter from Production Line	Sample C Floor Sweepings	Sample D Solid Paint waste with Filters & Paint
* * * * * * * * * * * * *	* * * * * *	* * * * *	* * * * *	* * * * * *
% Solids	NO FREE	100	100	100
Weight Raw Sample (g)	101.6	101.0	101.3	101.5
Filters Used	AP15	AP15	AP15	AP15
	HAWP	HAWP	HAWP	HAMP
* * * * * * * * * * * * * *	* * * * * *	* * * * * *	* * * * *	* * * * *
Initial Extract pH	9,4	5.3	8.8	6.1
Final Extract pH (24 hr)	5,6	5.3	5.5	5.9
Acid Added (24hr)(ml)	406.4	2./	112.2	36.
Final Extract pH (28hr)	муничну	5.1	5./	5.2
Acid Added (4hr)(ml)	<b></b>	2.	3/.	30.
Total Acid Added (ml)	406.4	4.1	143.2	66.
Total DI Water Added (ml)	1626.	20/6.	1883.	1964.
Original Liquid Phase(ml)	0	0	0	0
Final Extract Volume (m1)	2032.4	2020./	2026.2	2030.
* * * * * * * * * * * * *	* * * * * *	* * * * *	* * * * * *	* * * * *
RCRA Metals				
Arsenic (PPM)	<0.01	<0.01	0.023	<0.01
Barium (PPM)	<0.5	<0.5	1.0	1.1
Cadmium (PPM)	0.80	0.64	0,19	0./2
Chromium (PPM)	0.07	<0.05	<0.05	<0.05
Lead (PPM)	0.45	<0.2	<0.2	<0.2
Mercury (PPM)	<0.001	<0.001	<0,001	40.001
Selenium (PPM)	< 0,01	<0.01	<0.01	0.01
Silver (PPM)	<0.05	40.05	<0.05	<0.05
* * * * * * * * * * * * *	* * * * * *	* * * * *	* * * * * *	* * * * *

Note: Tables of pH adjustments with time have been extracted from the EIS sample work sheets and are reproduced in the section of this report containing laboratory bench sheets.

# ANALYTICAL REPORT SHEET SUPPLEMENTAL ANALYSIS - HAZARDOUS WASTES

This supplemental report sheet presents results of Corrosivity, Ingitability and Reactivity analysis if so required by the client. The following definitions apply to this analysis. The method of analysis is listed on the reverse side of this form.

- 1. Corrosivity A solid waste exhibits the characteristic of
  Corrosivity if it is aqueous and has a pH of less
  than (<) or equal to 2 or greater than (>) or
  equal to 12.5
- 2. Ignitability A solid waste exhibits the characteristic of ignitability if it is a liquid, other than an aqueous solution, containing less than 24% alcohol by volume, and has a Flash Point less than 140 °F (60 °C)
- 3. Reactivity Per procedures required by the Indiana State Board of Health, Reactivity is evaluated on the basis of Cyanide and Sulfide presence in the sample

Client Reactivity (ppm) \* Corrosivity Ignitability EIS Lab Sample Sulfide Cyanide (°F) (%C) Description (Hq) Number Sample A-Comp. Pretreatment Waste 6.4 > 158. 3.9 9.4 6516 Sample B-Paper 6.6. > 195. 3,9 51 6*5*2*G* Filter, Production NO FLASH-EXTINGUISHES 3.1 Sample C-Floor 5.8 9.1 6536 FLAME @ 140 Sweepings Sample D-Sold Paint waste 2.0 2.8 6.4 > 198 6546 200 North 200 Smet

\* Dry Weight Basis

#### METHOD OF ANALYSIS

The following procedural summaries are provided for clarification. The specific steps involved can be found in the following reference commonly referred to as SW-846: "Test Methods for Evaluating Solid Waste, Physical/Chemical Methods" USEPA, SW-846, July 1982, 2nd Edition.

#### CORROSIVITY (SW-846 Method 9040)

- . Liquid Waste (aqueous) are analyzed directly for pH using a commercial pH meter.
- . Solid wastes are analyzed for pH by preparing a 10% W/V solution of the waste in deionized water.

#### IGNITABILITY (SW-846 Method 1010 or 1020)

The method of analysis consists of heating a sample at a slow, constant rate with continual stirring in a closed cup tester. The Flash Point is defined as the lowest temperature at which a test flame applied to the sample ignites the vapors above the sample.

## REACTIVITY (SW-846 Method 9010 - Cyanide and Method 9030 - Sulfide)

- Cyanide is determined by refluxing of the sample under acid conditions with subsequent trapping of HCN gas in a Sodium Hydroxide scrubber. The scrubber contents are then analyzed either by titration or auto analyzer.
- . Sulfide is determined by sweeping Hydrogen Sulfide gas out of an acidified sample using a Nitrogen atmosphere. The HS gas is trapped in dual scrubbers containing Zinc Acetate and then analyzed by titration with Sodium Thiosulfate.

It is also possible to analyze a sample by the method of additions and to calculate the results, rather than plotting an analytical curve. The equation to be used for the calculation is shown below. The following three criteria must be met in order for the calculation to be valid:

- All solutions are made to the same final volume
- 2. A plot of concentration vs. absorbance is linear over the entire concentration range covered
- All readings are either in absorbance or in some value directly proportional to absorbance ("concentration")

$$C_{s} = \frac{C_{a} \cdot R_{s} \cdot V_{a}}{V_{s} (R_{a} - R_{s})}$$

where

C<sub>s</sub> = concentration of the element of interest
 in the original sample solution, in μg/ml
C<sub>a</sub> = concentration of the element of interest
 in the standard solution used to "spike"
 the diluted sample solution, in μg/ml
V<sub>s</sub> = volume of original sample solution used,
 in ml
V<sub>a</sub> = volume of standard solution used, in ml

 $R_s$  = reading obtained for diluted sample solution  $R_s$  = reading obtained for "spiked" sample solution

# THOD OF STANDARD ADDITIONS FOR METALS

EIS # 651G

TETAL Ag DATE	3-30-87 C	alibration 9	Std. 2 Abso	rbance .156	PLATE 100
Sample (m1)	Water (m1)	Std. (ml)	mg/l added	Result Cal	.c (<0.05) 0.0000 0.0136 0.0143 0.0141
3.50	1.50	Ø.ØØ	Ø.ØØ	.01	
3.50	1.00	.5Ø	.2Ø	.22	
3.50	.50	i.ØØ	.4Ø	.41	
3.50	0.00	i.5Ø	.6Ø	.62	
TETAL PB DATE	3-30-87 C	alibration S	Std. 5 Abso	rbance .046	
Sample (ml)	Water (ml)	Std. (m1)	mg/l added	Result Cal	0.0000
3.50	1.50	Ø.ØØ	Ø.00	.27	0.0000
3.50	1.00	.5Ø	.50	.66	.4792
3.50	.50	1.ØØ	1.00	1.09	.4617
3.50	0.00	1.5Ø	1.50	1.66	.4071
HETAL Cd DATE	3-20-87 C	alibration 9	Std. 2 Abso	rbance .221	PLATE 85
Sample (ml)	Water (ml)	Std. (m1)	mg/l added	Result Cal	.6 (0.80)
3.50	1.50	0.00	Ø. ØØ	.58	0.80)
3.50	1.00	.50	. 2Ø	.80	.7708
3.50	.50	1.00	. 4Ø	1.00	.7986
3.50	0.00	1.50	. 6Ø	1.19	.8217
ETAL Cr DATE	3-23-87 C	alibration S	8td. 5 Abso	rbance .167	
Sample (m1)	Water (ml)	Std. (ml)	mg/l added	Result Cal	.c
3.50	1.50	0.00	0.00	.04	
3.50	1.00	.50	.50	.45	
3.50	.50	1.00	1.00	.92	
3.50	0.00	1.50	1.50	1.36	
ETAL Ba DATE	3-23-87 C	alibration S	Std. 20 Abs	orbance .124	
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	Result Cal	0.0000
3.50	1.50	Ø.00	Ø.ØØ	.28	0.0000
3.50	1.00	.50	2.ØØ	2.34	0.3883
3.50	.50	1.00	4.ØØ	4.45	0.3837
3.50	0.00	1.50	6.ØØ	6.29	0.3993

#### HOD OF STANDARD ADDITIONS FOR METALS

IS # 6526

IETAL Ag I	DATE 3-30-87	Calibration S	Std. 2 Abso	rbance .156	
Sample (n 3.50 3.50 3.50 3.50	nl) Water (m 1.50 1.00 .50 0.00	0.00 .50 1.00	0. 00 . 20 . 40	Result Cal .01 .21 .41 .62	0.05 0.0000 .0143 .0143 .0141
IETAL Pb I	DATE 3-30-87	Calibration S	Std. 5 Abso	rbance .046	PLATE 102
Sample (n 3.50 3.50 3.50 3.50		0.00 .50 1.00	0.00 .50 1.00	Result Cal .05 .62 1.11 1.45	
1ETAL Cd I	DATE 3-20-87	Calibration S	Std. 2 Abso	rbance .221	
Sample (n 3.50 3.50 3.50 3.50	.50	0.00 .50 1.00	mg/l added Ø.ØØ .2Ø .4Ø .6Ø	Result Cal .46 .67 .87 1.08	0,64 0.0000 2,7%(CSO .6190 .6265 .6240
IETAL Cr I	DATE 3-23-87	Calibration (	Std. 5 Absc	orbance .167	
Sample (r 3.50 3.50 3.50 3.50		0.00 .50 .1.00	mg/l added Ø.ØØ .5Ø 1.ØØ 1.5Ø	Result Cal 0.00 .51 1.02 1.56	0.0000 0% RSD .0001 .0001 .0001
IETAL Ba	DATE 3-23-87	Calibration	8td. 20 Abs	sorbance .124	
Sample (1 3.50 3.50 3.50 3.50	ml) Water (n 1.50 1.00 .50 0.00	9. 00 1 .50 1 .00	mg/l added Ø.ØØ 2.ØØ 4.ØØ 6.ØØ	Result Cal .15 2.33 4.17 6.28	1c 0.0000 0%RSO .1966 .2132 .2097

### THOD OF STANDARD ADDITIONS FOR METALS

### :IS # 652GDUP

IETAL	Pb I	DATE	3-30-8	7 Ca	librat	ion	Std.	5	Absc	rbance.	Ø46	PLATE	102
Agri	3.50		1.	50 00 50	Ø. 1.	00 50 00	***	۷. 1.	00 50	Result .03 .53 i.08 i.46	•	0.0000 0.0000 0.354 0.340 0.373	
1ETAL	Cd I	DATE	3-20-8	7 Ca	librat	ion	Std.	2	Absc	orbance .	221	PLATE	85
	nple (n 3.50 3.50 3.50 3.50		1. 1.	. 50 . 00 . 50	Ø. 1.	00 50 00		Ø.	00 20 40	Result .46 .66 .87		0.0000 .6740 .6411	
IETAL	Cr I	DATE	3-23-8	7 Ca	librat	ion	Std.	5	Absc	rbance.	. 167	PLATE	92
	3.50 3.50		1,	. 50 . 00 . 50	Ø. i.	. ØØ . 5Ø		21. 1.	00 50 00	Result Ø. ØØ . 5Ø 1. Ø2 1. 56		0.0000 .0001 .0001	
Sai		ml)	Water 1	(ml) .50 .00 .50	Std. Ø.	(ml) .00 .50	) mg.	/1 0. 2.	added ØØ ØØ	Result .23 2.34 4.52 6.17	Cal	0.0000 .3114	94

## "HOD OF STANDARD ADDITIONS FOR METALS

:IS # 653G

IETAL Ag DATE	3-30-87 Ca	dibration Std.	2 Absor	bance .156	
Sample (ml)	Water (ml)	Std. (ml) mg	/l added	Result Cal	0.0000
3.50	1.50	Ø.ØØ	0.00	.01	0.0000
3.50	1.00	.5Ø	.20	.22	.0136
3.50	.50	1.ØØ	.40	.42	.0139
3.50	0.00	1.5Ø	.60	.58	.0150
IETAL Pb DATE	3-30-87 Ca	alibration Std.	5 Absor	bance .046	
Sample (m1)	Water (ml)	Std. (ml) mg	/l added	Result Cal	0.0000
3.50	1.50	Ø.ØØ	0.00	Ø. ØØ	0.0000
3.50	1.00	.50	.50	. 44	0002
3.50	.50	1.ØØ	1.00	. 97	0001
3.50	0.00	1.50	1.50	1. 45	0001
IETAL Cd DATE	3-20-87 Ca	alibration Std.	2 Absor	bance .221	PLATE 85
Sample (ml)	Water (ml)	Std. (ml) mg	/1 added	Result Cal	.c (0.19) 0.0000 .1951 .1951 .1905
3.50	1.50	Ø.ØØ	Ø.ØØ	. 14	
3.50	1.00	.5Ø	.2Ø	. 35	
3.50	.50	1.ØØ	.4Ø	. 55	
3.50	0.00	1.5Ø	.6Ø	. 77	
TETAL Cr DATE	3-23-87 Ca	alibration Std.	5 Absor	bance .167	PLATE 92
Sample (ml)	Water (ml)	Std. (ml) mg	/l added	Result Cal	0.05
3.50	1.50	0.00	0.00	.02	0.0000
3.50	1.00	.50	.50	.52	.0286
3.50	.50	1.00	1.00	1.05	.0277
3.50	0.00	1.50	1.50	1.56	.0278
METAL BA DATE	3-23-87 Ca	alibration Std.	20 Absc	rbance .124	PLATE 94
Sample (ml)	Water (ml)	Std. (ml) mg	/l added	Result Car	1c
3.50	1.50	0.00	0.00	.76	0.0000
3.50	1.00	.50	2.00	2.91	1.0100
3.50	.50	1.00	4.00	5.11	.9984
3.50	0.00	1.50	6.00	7.11	1.0259

#### HOD OF STANDARD ADDITIONS FOR METALS

:IS # 654G

IETOL On	NATE	マーマウータフ	Calibration	Std. 2 DE	aeocharce .	156 DLATE	100
~31							0.05
			.) Std. (ml)			Carc	0,03
3,50		1.50	0.00 .50			0.0000 .0301	
3.50 3.50		1.00	1.00				
3.54			1.50				
	<b></b> -						
IETAL Pb	DATE	3-30-87	Calibration	Std. 5 At	osorbance .	Ø46 PLATE	102
					1 25 .7.1		0,2
			l) Std. (ml)	***	ed Result	Ø.0000	
3.50 3.50		1.50 1.00				. 1111	
3.50		.50					
3.5		Ø. 00					
<b></b> .	_						
IFTOL Cd	DATE	3-20-87	Calibration	Std. 2 A	bsorbance .	221 PLATE	85
:	Air ( ) No.	tur Parritis to 1					0.12
Sample	(ml)	Water (m:	i) Std. (ml)		ed Result	Caic	Charles and Charle
3.5		1.50				ଡ. ଅଅଅଷ	
3.5		1.20				.1180	
3.5		. 50				. 1293 . 1253	
3, 5	Ø	ଅ. ଉତ	1.00	= (C) (C)	# / U	. ILUU	
TTAL Co.	NATE	70707	Calibration	Std 5 O	beorbance .	167 PLATE	92
ICIHL U	DM C	aEa01	MATTO! GOTAL!	0001 0 11	and the second s		0.05
Sample	(m1)	Water (m	l) Std. (ml	) mg/l add	ed Result	Calc	0.00
3.5	Ø	1.50			. Ø2	0.0000	
3.5	Ø	1.00	.50	. 50		. 0304	
3.5	Ø	. 50					
3.5	Ø	0.00	1.50	1.50	1.46	. 0298	
1							
'ETAL Ba	DATE	3-23-87	Calibration	Std. 20	Absorbance	.124 PLATE	
Sample	(ral)	Water (m	l) Std. (ml	) mp/l add	ed Result	Calc	(1.1)
3.5		1.50		0.00	.83	ଡ. ଡଡଡେଡ	
3.5		1.00	. 50	2.00	2.93	1.1293	
3.5	12)	. 50		4.00	5.00	1.1374	
3.5	iØ	ପ. ଅପ	1.50	6.00	7.17	1.1221	

## METHOD OF STANDARD ADDITIONS FOR METALS One Calibration Standard Used

Se 3-24-8	7 EIS #6	<i>516</i> Ca	libration Sto	1. 0.0200	(0.01)
Sample (ml)	Water (m1)   /2   9   6   3	Std. (ml)   O   3   6   9	0,004 0,004 0,008 0,002	Conc. (mg/l)   0.0002   0.00/8   0.0049	-3(4.9-,2) 2 2.005
Se 3-24-91	,			i. 0.0200	02(.2)9 = <.005
Sample (m1)	<del></del>	Std. (m1)		Conc. (mg/1)   0.0006	
Se 3-24-8	?7 EIS #	6536 Ca	libration St	d. <u>0,0200</u>	<0.01
Sample (m1)	Water (m1) /2	Std. (ml)     <u> </u>	mg/1 added 0.004 0.008 0.012	Conc. (mg/1)   <u>0.0008</u>   0.0029   0.0044   0.0056	$\begin{array}{c} .02(.8)3 \\ 3(2.98) \\ \hline .02(.8)6 \\ \hline 3(4.48) \\ .02(.8)9 \\ \hline 3(5.68) \\ \hline \end{array}$
	EIS #			d. <u>0,0200</u>	
Sample (ml)	Water (ml)    12   9   6   3	Std. (ml)   <u> </u>	mg/l added 0.004 0.008 0.012	Conc. (mg/1)   0.002   0.006   0.0/04   0.0/5	-3(6-2) -102(2)6 -3(10.4-2) -102(2)9, 10092
	EIS #	Ca	alibration St	.d	3(15-2)
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	Conc. (mg/1)	

# METHOD OF STANDARD ADDITIONS FOR METALS One Calibration Standard Used

	,				
As 3-24-87	EIS # 65	51 G Ca	libration Sto	1. 0.0200	(<0.01)
				Conc. (mg/l)   0.000/   0.0026	
Sample (ml)	Water (ml)	Std. (ml)	mg/l added	Conc. (mg/1)	.02(.1)4
4	<u></u>	<u> </u>		0.0001	4(2.6-1)
				0.0026	a2(d)8
	8	8	0,008	0.0069	102(.1)8 = 2.005
	4	/2	210.0		4(11.3-1) = <.005
					4(11.3-11)-2.00
And the second s					
As 2-24-87	EIS # 6.	52.6 c-	libration St	0.0200	
713 32707	_ ETD #	ta	TITDESCION SO	1	· (<0,01)
State - Company	Tilledonia (m.1.)	CF4 (m1)	ma/l added	Conc. (mg/l)	
Sample (ml)	Water (ml)   /6	O I	mq/1 added	0.0014	4(4.2-1.4) .010
	/2	4	0.004	2 22/2	4(4,2-1.4)
	8	8	0,008	0.0098	-,02(1.4)B -4(9.8-1.4)=.007
	4	/2	0,0/2	0.0137	
	7	121		1	02(1.4)12 .007 4(13.7-1.4)
•					4(13.7-1.4)
٠.					
As 3-24-8	37 mm 4	536 c	alibration St	d. <u>0,0200</u>	
	. FID #	<del>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</del>	XIIDEGCION DO		- (0.023)
Cample (m1)	Nator (ml)	Std (m1)	l ma/l added	Conc. (mg/1)	
Sample (mil)	Wares Imil	Ded. (mr)	1 0	0.0032	- 102(3.2)4025
<u> </u>	/2	4	0.004		
	8	8	0.008	0.0088	-, <u>02(3,2)8</u> -4(8.8-3,2)
		12	0.0/2	1 /	•
<u> </u>	<u> </u>		1	1 0101-1	02(3.2)12 , .022 4(12.1-3.2)
					4(12.1-5.4)
4					
As 2.26.	<i>-87</i> EIS #	654 G C	alibration St	d. 0.0200	
113 5-64	-, FT9 #		MIIDIMOIO. DO		- (<0.01)
: :) - <b>::</b>	1 Marketon (m.1)	i C+3 (m1)	l ma/1 added	Conc. (mg/1)	
Sample (MI)	Water (MI)	Sca. (mil/	1 0	0.00/2	- 102(1,2)4
4	<u> </u>	4	0.004	0.0051	- 4(S.1-1.2) · .006
	/2	8	0.008	0.0106	02(1.2)8 - 4(10.6-1.2) ,005
·	8	12	0.012	0.0144	
<u> </u>	1	1			02(1.2)12 4(14.4-1.2)
					4(144-119
	ETC	C	alibration St	- d	
<u></u>	EIS #		althracton b		<del>_</del>
		. CL3 /11	I ma/1 nddad	Conc. (mg/1)	
Sample (ml)	Water (ml)	Std. (ml)	i md/i added	1 COIC MAY 17	
			. ]	]	
		1	<u> </u>	_ [	
			.   <u></u>	_	<del></del>
	i	1	1		

## Mercury Worksheet

c	ulfuric Acid (H2S	na) 5 3-	Pot. Dichro	omate (K2Cr2O7)							
N	itric Acid (HNO3)	2,5 3.	<b></b>	ydrox. Hydrochloride							
			0000 0-12								
	ot. Permanganate			1 NH2OH·HC1)							
F	Pot. Persulfate (K2S2O8) 8 3-9 Stannous Chloride (SnCl2) 5 3-2										
•	bring all up to 150 ml										
	Bottle #	Sample #	Vol. or Wt.	Result							
L	(	131		heightum Z							
2	7	731		2.5							
3	14	0.05 ug Ha		3							
4	17	0.1 119		13							
5	<u></u>	0.2 1/4		24							
6	18	0.3 7.0		37							
7	21	0.4 119		50							
8	23	1110	100ml	2.5							
9	78		100 to. 1/19 Ha	/3							
	82	<del> </del>		27							
0	84		100 +0.2/19	37							
1	98	120(6)	100 to.3 deg	3							
2	100	395©	100-								
3, :	101		100 to. long Ha	28 20,001							
4	107		100 40.2/19								
5	116		100 + 0,3/19	44							
6	117	396 ©	100ml	2.5							
7	118		100 tollaffa	15 (20.00)							
8	129		100 40.2 119	29							
^	141		100+0.3 Tea	42							
0	144	wp016-2	Zonf	35 0,01533 102,296,6							
		act = 0.015	<del></del>								

COMMENTS

#### Mercury Worksheet

Sulfur	cic Acid (H2SC	)4) <u>5</u>	
Nitrio	e Acid (HNO3)	2.5	
Pot. F	Permanganate	(KMnO4) <u>5</u>	
Pot. F	ersulfate (K	2s208) <u>වි</u>	

COMMENTS

Pot. Dichromate (K2Cr2O7)

Sod. Cl.-Hydrox. Hydrochloride

(NaCl NH2OH·HCl) 5

Stannous Chloride (SnCl2) 5

- Marie Marie	Bottle #	Sample #	Vol. or Wt.	Result
- maken	171	3970	100m/	height mm
: 3	176	(	100 to. King Ha	/3
	179		10640.2/4	26 20.00 250
	182		100+0.379	38
,	183	3976 Dup	Toom	4.
;	185	1	100 +0. Kg Hg	14 (20.001)
,	191		10010.2/14	28
}			10010.34	43
,	230	5950	100ml	4
)	232	1	100 +0. (seg Ha	
	233	<del> </del>	100 +0.2/19	3º vr/
}	238	1	100 +0.3/19	
3	241	618C)	100 m	3.5
1	244	+ (	100ml +0.1 eig the	16
ng pychy a	250		1, 1, 1, 1, 1,	29 (20.00)
5	254		loom to Eleg	42
5	[ ]	651 (0)	100 ml + 0.3dg	5
7	278	1	100ml Hz	
3	363	<del>                                     </del>	100 + 0.1 119	27 (20:00)
) 	365		100 40 219	39
َ	3 79		100+0.3 leg	

#### Mercury Worksheet

Sulfu	ric Acid (H2SO4)	5	<u> 3-2</u>
pro man	ic Acid (HNO3)		<u> 3-</u> 2
	Permanganate (KM		<u>3 -</u> 30
	Persulfate (K2S2	- 22	3-3

	Bottle #	Sample #	Vol. or Wt.	Result
1	8	131		height mm
2	9	BI		O
3	I	0.05 ug Ha		4
4	/.3	0.1 119		3.5
5	16	0.2. 119		17
6	22	0,3/19		26
7	24	0.4 119		36
8	29	5956 0	2~1	6
9	31		2ml +0./1/9 Hg	12.5
0	32		2ml +0.2 gra	24 0.02
1.	3.5		2ml +0.3 pg	36
2	37	6526 C	100 m/	0.5
3	44		100 ml to lugty	10 (=0.001)
4	45		100 ml +0. 200 Hig	18
5	48		100 m/ 10,300	24
6	74	6536 ©	100ml	1.5
7	83		poul + o. las Ha	"
.8	86		100 / to. 200g	22 (20.00)
9	88	/	100m/10, mg	34
U	163	wp016.2	20	32 0.0181 120,8% Rac

act = 0.015 95% c.I. = 0.011 - 0.0194

COMMENTS

<u>3</u>.24

3.18

## Mercury Worksheet

Sulfuric Acid (H2SO4) 5 3-1	Pot. Dichromate (K2Cr2O7)
Nitric Acid (HNO3) 3.5 3-2	Sod. ClHydrox. Hydrochloride
Pot. Permanganate (KMnO4) 5 3-30	(NaCl NH2OH·HCl) 5
Pot. Persulfate (K2S2O8) 8 3-)	Stannous Chloride (SnCl2) 5

	1		Result
Bottle #	Sample #	Vol. or Wt.	Result
112	6546 D	100 ml	height min
113	·	100 ml + 0.1 sig Hg 100 ml + 0.3 sig	12 20.001
114		100-1+0,2/19	22
127		100 ml + 0.3 leg	15.
133	748 G (C)	100.7	
138	. (	100-lto. 1 sight	12
169	(	100 1 to. 2 seg.	21.5 (20,001)
		100 1 to.3 lea	35
174	7496 O	100mt	
	1		11 (0.001)
188		100ml to. 27/4 100ml to. 27/4	23 0% RSD
	1	100-110.3110	36
217	7496 (D)	100ml	٥
222	(	100. 1 + 0. 1 x19 Hg	12
231		100. 10. 2jig	24 (20.001)
263	1	100.01 10.364	37
363	750 6 W	100,0	0
361		100ml + 0. king Hay	12 (20,001)
372			25
3 78		100m/ 10,3119	40
379	1	1 room respect	

COMMENTS

## SOLID SAMPLE CYANIDE WORK SHEET

								Anal	ysis Informa	tion	
		Not No	eight Informa	tion		%	Sulfide	Scrubber	m1	Auto	
	EIS Lab	wet we	eight informa		Liquidy		in	Vol	Titrant	Analyzer	Cyanide (ppm)*
Date }	Number	Final	Initial	Net	Volume	M	Scrubber	Titrated	(0.00282N)	(mg/1)	(ppiii) *
4w	Blank				500		NP	50	0.15		
3-5-87	4716-8	75.3	65.3	10.0	కరాల	.45	4N_	50	0.2		1.48 *
	<b>6</b>										*
	6180	85.0	74.3	10.7	500	60.	NP	50	0.35		13.7*
	6	•									00*
	6510	93.4	71.0	12.4	500	69.6	NP	50	0.2		3.9*
	0										111108*
$\underline{\checkmark}$	17/0	76.6	66.3	10.3	500	104	NP	50	0.15		< 4.28*
иш											
رة - اع 12-	Blank				500	<u> </u>	NP	50	0.1		
\ \ \ /	9					1					
	6536	112.1	100.8	11.3	500	9.92	NP	50	0.3		5.77*
	(1)			<u> </u>	1						*
	6546	110.8	0.00	10.8	500	2.16	NP	50	0.2		2.78*
	(3)					1					
	6526	107.7	100.0	7.7	800	1.95	NP	50	0.2		3.89*
											Pac 94.0%
~	5.0-4/450				500		NP	50	7.3		4.2mg/L
Uju											
3-19-87	1				500		NP	50	0.1		
3-11-61	1316										*
	6976	75.7_	65.7	10.0	55-0	•02	NP	50	0.15		1.47*
	3				-						1.45*
$\forall$	6986	75.0	64.9	10.1	500	-O3	NP	50	0.15		1.73

#### PENSKY MARTENS CLOSED CUP FLASH POINT LOG

Analyst 4w Analysis Date 2-27-87

#### P-XYLENE READINGS

Trial #	Observed Value C F	Barometric Pressure (mmHg)	Corrected Flash Point C F	Average Value Corrected Flash	Quality Control Limit
1	81°	740	<b>8</b> 2.ລ <sub>ີ</sub>		26.2 - 27.8 Centigrade
2	7 <del>9°</del>	740	80.2°	81.2°	79.2 - 82.5 Fahrenheit

#### SAMPLE ANALYSIS

#	Sample Type L=Liquid S=Solid	Observed Value C F	Barometric Pressure (mm Hg)	Corrected Flash Point C F	Comments and/or Precision data
6510	S	>157°	740	>158.2°	
6970	S	>198°	740	>199,2°	RSD
697 DUP	S	>197	740	>198.2°	0.4 %
65.20	3	> 194°	740	>195.2°	
6546	S	> 197°	740	>198.2°	orange flome ad 75°
	-				
			·	•	
			,		

#### CALCULATION FOR CORRECTED FLASH POINT

Analysis :	Date	3-26-87	Analyst	MW

#### P-XYLENE READINGS

	Trial #	Observed Value C	Barometric Pressure (mmHg)	Corrected Flash Point C (F)	Average Value Corrected Flash C F	Quality Control Limit
	1	79°	738	%0.3 ఄ		26.2 - 27.8 Centigrade
_	2	80°	)3 <i>&amp;</i>	81.3	30.9°	79.2 - 82.5 Fahrenheit

#### SAMPLE ANALYSIS

Sample Type L=Liquid S=Solid	Observed Value C F	Barometric Pressure (mm Hg)	Corrected Flash Point C F	Comments and/or Precision data
S		738		red he at 101° TAKEN TO 1560
				•
			_	
	L=Liquid S=Solid	L=Liquid Value S=Solid C F	S=Solid C (F) (mm Hg)	L=Liquid Value Pressure Flash Point S=Solid C F (mm Hg) C F

#### CALCULATION FOR CORRECTED FLASH POINT

- Using oF and mm Hg CORRECTED FLASH = (Observed Value) + [0.06 (760 Barometric Pressure)]
   Using oC and mm Hg CORRECTED FLASH = (Observed Value) + [0.033 (760 Barometric Pressure)]

#### PH OF SOLID SAMPLES & CORROSIVITY

				SAM 1	SAMPLE APPEARANCE			pH MEASUREMENT MADE ON 10% OTHER		
DATE	ANALYST	TIME	EIS#	LIQ.	LIQ.	SOLID	LIQ.	W/V SOLUTION	(SPECIFY)	pН
2-11-87	CB	1:53	3956			/	-			9.2
	Ì	1:57	396			V				9.4
		2:02	397			V				9.5
2-13-87	CB	10:48	4266			<i>i</i>		2	and the second s	12.
· 1	CB	10:53	4276			<u></u>			-	10.8
	DUPLICATE	ANALYSIS				1				10.8
2-16-87	CB	3:18	4516			~		<u></u>		6./
2-18-87		10:38	6							8.8
3-6-87		7:15	6186			-		U		7./
3 / 1		9:33	6516							9.4=
V	1	9:20						2-	•	5./

## SPECIAL INSTRUCTIONS FOR CORROSIVITY SAMPLES

- 1. If sample is liquid or semi-liquid, pH is taken on the as is sample.
- 2. If sample is a solid, weigh out 10 grams, place it in a 150 ml. beaker and bring to the 100 ml. beaker mark with DI water
  - Stir the suspension several times during the next hour with a glass stirring rod
  - Sample is placed on a magnetic stirrer and gently stirred while measuring the pH
  - Analysis is complete when the sample has attained a relatively stable reading where pH does not change by more than 0.1 units in 30 seconds

## PH OF SOLID SAMPLES & CORROSIVITY

				SAMPLE APPEARANCE			pH MEASUREMENT MADE ON 10% OTHER			
DATE	ANALYST	TIME	EIS#	LIQ.	LIQ.	SOLID	LIQ.	W/V SOLUTION	(SPECIFY)	рН
3-6-87	CB	9:37	653°			~				9.1
4	<b>V</b>	9:25	.6							6.4
3-14-8	7 CB	11:30	1716			V				7,9
	CB	12:01	7406							9.5
	CB	11:55	1-6							8.9
	DUPLICATE	ANALYSIS	7486					V		8.8
	CB	11:40	6966					V		7.6
		11:43	6976			/		V		7.4
		11:46	6986			V		V ,	·	7.4
		11:50	6980			V		V		7.3
	V	11:57	7506							8.6

### SPECIAL INSTRUCTIONS FOR CORROSIVITY SAMPLES

- 1. If sample is liquid or semi-liquid, pH is taken on the as is sample.
- 2. If sample is a solid, weigh out 10 grams, place it in a 150 ml. beaker and bring to the 100 ml. beaker mark with DI water
  - Stir the suspension several times during the next hour with a glass stirring rod
  - Sample is placed on a magnetic stirrer and gently stirred while measuring the pH
  - Analysis is complete when the sample has attained a relatively stable reading where pH does not change by more than 0.1 units in 30 seconds

## PH METER STANDARDIZATION

			1	BUFFER	USED	,	
NALYST	DATE	TIME	TEMP.	ACTUAL / READS	ACTUAL / READS		QC
<u>GB</u>	2-18-87	3:32	220	7.41/7.42	4.01/4.01	 <u></u>	
Sto	2-19-87	9:02	210	7,41	10.4/10.39		 
CB	2-19-87	11:27	220	7.41/7.40	4.61/4.01	 	
· CB	2-20-87	2:52	220	7.41/7.41	4.01/4.01		
JW	2.23.87			7.41/7.42	4.01/10,39	 · · · · · · · · · · · · · · · · · · ·	
M	2-2487	10:14	210	7.41/7.40.	10,4/		<u> </u>
CB	2.24-87		210	7.41/7.46	4.01/4.01		ļ
C/S_	2-25-87			7.4//7,40	4.01 (4.01		
IW	2-27-87	į.	220	7.41/7.41	4.01/ 4.01		
IN	3-2-87			7.41/7,43	10,4/10,40		
CB			20°	7.41/7.4/	4.01/4.00		<u> </u>
CB	3-3-87		•	7.41/7.41	40//401		-
CB.	3-4-87			7.41/7.40	4.01 / 4.0 (		
B	3-5-87		248	7.4//7.41	4.01/4.01		
CB	3-6-87	200	220	7.41/7.42	4.01/4.01		-
CA	3-6-87		220	7.4// 7.41	10.4/10.40	<u></u>	
JW	3-6-87	10:09	210	7.41/7.41	4.01/4.01		
CB	3-7-87		220	7.41/7.41	4.01/4.00		
DU	3-8-87	10:45		7.41/7.41	4.01/4.01		
\	3-9-87	10:51	226	7.41/7.41	4.01/4.00	<u> </u>	
_CB	3-10-87		ا	7.41/7.43	10,40/10,39	,	

#### SULFIDE ANALYSIS WORK SHEET

	-									T	<del> </del>	
	1	Solid Sample Preparation				Analysis Information				Normal		
						Solid .	Liquid					ppm
	EIS Lab	Final	Initial	Net	Volume	Aliquot हैं (ml)(g)	Sample	Iodine	Titrant	<u>Iodine</u>	μg	mg/l
Nate	Number	WT (g)	. Wt (g)	Wt (g)	. DI (m1)	(ml)(g) 🗞	(ml)	(m)	(m1)	Titrant	<u> </u>	S
			ļ		1					0.0245	_	•
	ا ، ما				300			10.0	9.8	0,0250	0	
3-10-87	B				300		_ <del> </del>	70.0	1,0	0,0250		
	1											
	7066				300			10.0	9.6		80	0.26
	700					<del></del>						
					1	4					_	- 4
•	1716	2424	201.2	41.2		0		10.0	9.6		.80	2.0*
		0.100.1				<del> </del>				0.0245		
									200	<del></del>	0	
3-11-87	Bl		1		300			10.0	9.8	0.0250		
7 11												
				, ,		62.6		/0 -	9.6		80	6.4*
	6516	247.0	206.3	40.7		jõ		10.0	1.6		00	6.7
	/ <del></del>			-				į	1	1		
. /	1,> 6	2021		10 1		887	1	10,0	01		80	6.6*
1	6526	222.4	210.0	12.4		\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		10,0	9.6		00	0.0
						1 1	1	1.				
	653 G	208,6	168.5	40.1	1	1		10.0	9.5		112	3.1*
a de la constanta de la consta	653	208.6	168.5	70,1				10.0		•	,	- ·
								•			0 -	
	6546	241.7	201.5	40.2		<b>b</b>		10.0	9.6		80	2.0
	(6-)	271.1	201.5	10.2			<u> </u>	70.0	+			
												0.77 * \$\overline{x} = 3.4 4.92.850 4.0 *
:	1.966	209.3	11.78	41.5		(6		10.0	9.7		32	0.77*
	0.0	2011	167.0				<del>                                     </del>	' '	· · · · · · · · · · · · · · · · · · ·		!	¥=3.4
					1		1		9.4			14.92.250
	16976	247.8	217.4	40.4				10.0	7.7	1	160	4.0*
i	<del>  • •</del>	100	00111	<u> </u>	<del>- </del>	-		<u> </u>				į
•	200					į	ν				00	1
	698	208.4	168,6	40.0		1	<b>'</b>	10,0	9.6		80	2.0*
		+	, , , , ,	<del>                                     </del>	<del>                                     </del>							1.
	. 101		2/2 2	111			2)	122	9.6		80	1.9*
-3-16-87	16446	251.6	210.0	41.7		•		10.0	1,0		<u> </u>	1111
7 10 4			<u> </u>							i		

\* ppm dry

## HAZARDOUS WASTE FILTRATION

•	Date 3-5/3-6-87 Analyst <u>CB</u> EIS# 65/
2.	Sample & Date Vice Sinc 2-12-87
.3٠	Description of Sample: Sample A Composito of Photreaument Waste
4.	Type of Analysis: Metals Organics Other
5.	Wt. of raw sample used for first separation (100 gm. minimum) 101.6
6.	Filters Used Weight (gm.) Filters Used Weight (gm.) 211.9
	<u>HAWP</u> 110.3
	AP15 101.6
	NO FREE Liquid
7.	TOTAL WEIGHT +
8.	Total weight of solid material and filters after filtration
9.	- Less tare weight of filters used (line 7)
LO.	= Actual weight of material for extraction
11.	If the solid residue on the filters is less than 0.5% of the original
	weight of the waste, the residue can be discarded and treat the liquid  Result from line 10 ( ) $\times$ 100 =
	phase as the extract. Weight from line 3 (
12.	If the result from the formula in 11 is greater than 0.5% solids, you
	must go. through extraction. The total amount of distilled water to be used
	in transferring material to the extractor and which is to be used in the
	extraction is obtained by the following formula:
	Weight from line 10 ( $10/.6$ ) x 16 = $1626$
13.	The total amount of 0.5N Acetic acid which can be used in pH adjustment
٠	is obtained by the following formula:
7	Weight from line 10 (161.6 ) $\times$ 4 = 406.4
14.	With residue and distilled water in extraction container (92000 man)
	used if organics are to be analysed - line 4 ) agitate and measure pH.
	If pH is greater than 5.0 decrease to 5.0 $\frac{+}{2}$ 0.2 by adding 0.5N acetic acid

l.

## HAZ DOUS WASTE FILTRATION

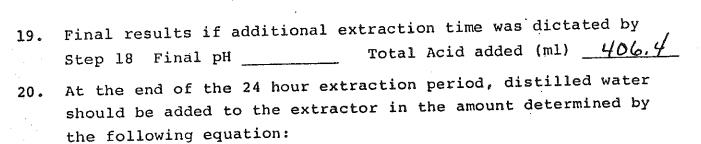
singled at 15, 30 and 60 minute
The pH of the solution should be adjusted at 15, 30 and 60 minute
The phron of the phrone interval if the phrone not have
The pH of the solution of the pH does not have intervals, moving to the next longer intervals, moving to the next longer interval if to $5.0^{\pm}0.2$ . This
than 0.5 pH units to bring it to 500
adjustment Must be carried on for AT LEAST 6 HOURS. This time will be
adjustment Must be carried on for AT LEAST & HOURS
haming by mole clian var i
adjustment Must be carried and adjust more than 0.5 pH units at longer if the sample is still changing by more than 0.5 pH units at 100 minute or less time intervals. Adjust pH only when it rises above 5.2
time intervals. Adjust ph only
60 minute or less time intervals. Adjusted pH Acid Added (ml.) 406.4
Adjusted pil

00 11121111			- 11 - Had DH	Acid Added (MIL.)	- 1
Clock time		рн 7, 4	Adjusted pH	130.0	é
8:37		8.1	5.0	200.0 330.0	
9:09		5.8	5.2	76.4 406.4	
_1.5/					
	**************************************				
		. <u></u> .			
,					
					•
				1ded 406.4	-
	- of A19:45 Final pH	<u>ک</u> ړي	Total Acid Ac	Ided	

17. If at the end of the 24hr. extraction period the pH of the solution is not below 5.2 and the maximum amt. of acid has not been added (line 13) the pH should be adjusted to  $5.0^{+}$  0.2 and the extraction continued for an additional FOUR hours, during which the pH will be adjusted at one hour intervals.

Clock Time	Time Interval	рН	Adjusted pH	ACIU Accus

## HAZARDOUS WASTE FILTRATION



$$V = [(20) (W) - 16 (W)] - A$$

$$V = [(20) (DLG) - 16 (10/G)] - (406.4) = 0$$

V = ml distilled water to be added

W = weight in grams of solid charged to extractor (line 10)

A = ml of 0.5N acetic acid added during extraction (line 17 or 19)

- 21. This liquid in the extractor is now filtered once again and the liquid obtained is combined with the liquid which was obtained in the first filtration. This combined sample is now ready for analysis.
- 22. Volume (ml) of liquid from first filtration

  Total Volume (ml) of DI water added to sample /626

  Total Volume (ml) of acid added to sample (Line 19) 406.4

23. COMMENTS:

24. Solid Sample Preparation

- $\sqrt{\phantom{a}}$  1. Completely sifted through a 9.5 mm standard sieve
- 2. A surface area per gram of material equal to or greater than 3.1 square centimeters
- 3. Been cut, crushed, or ground to the point where it may pass through a 9.5 mm standard sieve
  - 4. Been subjected to the Structural Integrity Procedure 3.

## HAZARDOUS WASTE FILTRATION

**.	Date 3-5/3-6-87 Analyst CB EIS# 652
	Sample & Date Viron Onc. 2-12-87
	Description of Sample: Sample B Paper Filter (Production Line)
9	Type of Analysis: Metals Organics Other
	Wt. of raw sample used for first separation (100 gm. minimum) /01,0
	Filters Used Weight (gm.) Filters Used Weight (gm.) 204.2
	HAWP 103.2
	AP15- 101.0
95- 14	
	DRY
,	TOTAL WEIGHT + =
} •	Total weight of solid material and filters after filtration
}	- Less tare weight of filters used (line 7)
)	= Actual weight of material for extraction
. •	If the solid residue on the filters is less than 0.5% of the original
, ,	weight of the waste, the residue can be discarded and treat the liquid  Result from line 10 ( X 100 =)  Weight from line 5 ()
,	If the result from the formula in 11 is greater than 0.5% solids, you
	must go. through extraction. The total amount of distilled water to be used
	in transferring material to the extractor and which is to be used in the
	extraction is obtained by the following formula:
	Weight from line 10 $(10/.0)$ x 16 = $16/6$
	The total amount of 0.5N Acetic acid which can be used in pH adjustment
	is obtained by the following formula:
	Weight from line 10 ( $101.0$ ) x 4 = $404.0$
•	With residue and distilled water in extraction container (glass Must be
, a	used if organics are to be analysed - line 4 ) agitate and measure pH.
	If pH is greater than 5.0 decrease to 5.0 $\frac{+}{2}$ 0.2 by adding 0.5N acetic acid.

#### HAZA JOUS WASTE FILTRATION

pH of the solution should be adjusted at 15, 30 and 60 minute
intervals, moving to the next longer interval if the pH'does not have
to be adjusted more than 0.5 pH units to bring it to $5.0\pm0.2$ . This
adjustment Must be carried on for AT LEAST 6 HOURS. This time will be
longer if the sample is still changing by more than 0.5 pH units at
60 minute or less time intervals. Adjust pH only when it rises above 5.2
404.0

Clock time	Time Interval	_рн_	Adjusted pH	Acid Added (ml.)
8:37	0.25	<u>5</u> ,3	5.0	2./
9:09	0.25	5,2		
9:37	0.50	5.2		
10:15		5.2		
11:25		5.2		
12:30	1.0	5.2		
1:34	1.0	5.2		
2:38	1,8	6.2		
3:45	18.0	5.2		
		<u></u>		
	off At 9:48			0 /
	Final pH	' <u>5,3</u>	Total Acid Add	$\frac{\partial}{\partial x} = \frac{\partial}{\partial x} $

If at the end of the 24hr. extraction period the pH of the solution is not below 5.2 and the maximum amt. of acid has not been added (line 13) the pH should be adjusted to  $5.0^{\pm}$  0.2 and the extraction continued for an additional <u>FOUR</u> hours, during which the pH will be adjusted at one hour intervals.

Clock Time	Time Interval	рН_	Adjusted pH	Acid Added (ml.)	900
9:55	1.0	<u>5,3</u>	<u> </u>	2.0 4.1	399.
10:58		5.0			
12:05	1.0	5,1			
1	1 10	ا سے			

## HAZARDOUS WASTE FILTRATION

~ 1245-41111

19.	Final results if additional extraction time was dictated by Step 18 Final pH $\frac{5.7}{2}$ Total Acid added (ml) $\frac{4.7}{2}$
20.	At the end of the 24 hour extraction period, distilled water should be added to the extractor in the amount determined by the following equation:
	$V = [(20) (W) - 16 (W)] - \Lambda$
· -	V = [(20)(101.0) - 16(101.0)] - (4.1) = 400
	V = ml distilled water to be added
	W = weight in grams of solid charged to extractor (line 10)
	A = ml of 0.5N acetic acid added during extraction (line 17
. Villa i	or 19)
21.	This liquid in the extractor is now filtered once again and the liquid obtained is combined with the liquid which was obtained
la. Parije	in the first filtration. This combined sample is now ready for
	analysis.
22.	Volume (ml) of liquid from first filtration  Total Volume (ml) of DI water added to sample 20/6  Total Volume (ml) of acid added to sample (Line 19) 4//
23.	COMMENTS:
24.	Solid Sample Preparation  1. Completely sifted through a 9.5 mm standard sieve
	2. A surface area per gram of material equal to or greater than 3.1 square centimeters
₩. ₩.	3. Been cut, crushed, or ground to the point where it may pass through a 9.5 mm standard sieve
	4. Been subjected to the Structural Integrity Procedure 3.
1.0	

#### VITCO. INC.

#### P. O. BOX 407, NAPPANEE, INDIANA 46550

PHONE; 219-773-3181

February 27, 1987

Mr. H. Stephen Nye, P.E. EIS Environmental Engineer, Inc. 1701 North Ironwood Drive South Bend, Indiana 46635

Dear Steve,

dia :

I am sending you four samples of waste streams for analysis of RCRA characteristics of ignitability, corrosivity, reactivity, and EP toxicity. The four samples were collected on February 12, 1987, and are labeled as follows:

Sample A: Composite of Pretreatment Waste

Sample B: Paper Filters from Production Line

Sample C: Floor Sweepings (Sawdust & P.E. Overspray)

Sample D: Solid Paint Waste with Filters & Paint Overspray

Samples A,B,C may have levels of various metals and Sample D may have levels of solvents, particularly xylol.

I am having these samples analyzed by request of Jeff Blankenberger, an inspector for the State of Indiana. I would appreciate the results as soon as possible. We had discussed a fee of \$510.00 per sample over the phone. I have a purchase order (number 84607) assigned to this order.

Please call if you have any questions or need any additional information.

Sincerely,

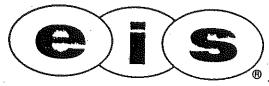
Vitco. Inc.

Brett L. Rordmann

Brett L. Nordmann Ceramic Engineer

BLN/nkn

cc: Robert Koehl, Vitco, Inc.



## EIS ENVIRONMENTAL ENGINEERS, INC.

1701 North Ironwood Drive • South Bend, Indiana 46635 • 219/277-5715

#### WASTE CLASSIFICATION ANALYSIS REPORT

erio. G				
Client: Vitco, Inc.	Sample Description			
ATTN: Brett L. Nordmann Date Sampled: Date Received: 3-2-87 Date Forwarded: 4-3-87 Purchase Order: 84607	EIS Analysis No.: 651G - 654G  651G - Sample A; Composite of Pretreatment Waste 652G - Sample B; Paper Filter from Production Line 653G - Sample C; Floor Sweepings 654G - Sample D; Solid Paint Waste with Filters & Paint Overspray			
No. 1				

This report presents results of waste classification through laboratory analysis procedures. The following references were utilized, as needed, in the evaluation procedures herein.

- "Test Methods for the Evaluation of Solid Waste Physical/Chemical Methods" USEPA SW-846, July 1982, 2nd Edition
- "Methods for Chemical Analysis of Water and Wastes" EPA 600/4-79-020
- State of Indiana "Leaching Method"

The specific client requested analysis for the samples described above were the following.

EP Toxicity - Metals	X	State of Indiana Leaching Method	
EP Toxicity - Organics		Volatile Organic Compounds	
Ignitability	X	Semi-volatile Organic Compounds	
Corrosivity	X	(Base/Neutrals Acid Fraction)	·
Reactivity	X	PCB Pesticides	
Additional			·

Materials constituting this report packet include laboratory analysis bench sheets. These bench sheets are required by the State of Indiana as an integral part of the Waste Classification Analysis Report. Certain sections of this report may not pertain to your samples but do constitute a part of the EIS Report Packet. All results are hand entered to eliminate data transfer errors.

Andris Rozite, Laboratory Director

# QUALITY ASSURANCE DATA EP TOXICITY and/or LEACHING METHOD

	% Recovery	% RSD		
Parameter	USEPA EMSL		Precision	
	QC Sample	Matrix Spike	Analysis	
Arsenic *	95.7	-		
Barium *	108.8	-	0 (6526)	
Cadmium *	108.	-	2.7 (6526)	
Chromium *	99.8	-	0 (6526)	
Copper				
Iron				
Lead *	94.	-	0 (6526)	
Manganese				
Mercury *	102.2, 120.8	_		
Nickel				
Selenium *	1/3.8			
Silver *	104.8	_		
Sodium				
Zinc	·			
Chlorides				
Cyanide, Total				
Fluoride				
PCB				
рН				
Phenols				
Sulfate				
Sulfide, Total				
TDS				
TOC				
тон				

<sup>\*</sup> These metals are analyzed by the Method of Standard Additions

## REGULATORY LIMITS (ppm)

To the second se	EP Toxicity		State of Indiana Leaching Method Foundry Waste Types Only			
Parameter	RCRA	A	B B	- C	D_	
Arsenic *	5.0	0.05	0.5	1.25	5.0	
Barium *	100	1.	10.	25.	100.	
Cadmium *	1.0	0.01	0.1	0.25	1.0	
Chromium *	5.0	0.05	0.5	1.25	5.0	
Lead *	5.0	0.05	0.5	1.25	5.0	
Mercury *	0.2	0.002	0.02	0.05	0.2	
Selenium *	1.0	0.01	0.1	0.25	1.0	
Silver *	5.0	0.05	0.5	1.25	5.0	
Chlorides	-	250.	2500.	6250.	* *	
Copper	-	0.25	2.5	6.25	**	
Cyanide, Total		0.2	2.	5.	**	
Fluoride	wate	1.4	14.	35.	**	
İron	<b></b>	1.5	15.	**	**	
Manganese	<b></b>	0.05	0.5	**	**	
Nickel	_	0.2	2.	5.	**	
PCB	-	_		_	_	
рн	<b></b>	6.0 - 9.0	5 - 10	4 - 11	**	
Phenols	<del>-</del>	0.3	3.	7.5	**	
   Sodium		250.	2500.	6250.	**	
Sulfate	_	250.	2500.	6250.	* *	
Sulfide, Total		_	5.	12.5	**	
ŢDS	_	500.	5000.	12500.	* *	
TOC		<u></u>		-	_	
тон	_	ans	, <del>-</del>	_		
Zinc	<del></del> •	2.5	25.	62.5	* *	

<sup>\*</sup> Limits shown are based on EP Toxicity Analysis Data

<sup>\*\*</sup> Testing is not required

## ANALYTICAL REPORT SHEET EP TOXICITY - METALS ANALYSIS

			1	
EIS Lab Number	6516	6526	6536	6546
Client Description	Sample A Compose of Pratreatment Waste	Sample B Paper Filter from Production Line		Sample D Solid Pant waste with Filters & Pain Overspray
* * * * * * * * * * * * *		* * * * * *	* * * * *	* * ***
% Solids	NO FREE	100	100	100
Weight Raw Sample (g)	101.6	101.0	101.3	101.5
Filters Used	AP15	AP15	AP15	AP15
	HAWP	HAWP	HAWP	HANP
		•		
* * * * * * * * * * * * *	,	* * * * *	* * * * * *	* * * * *
Initial Extract pH	9.4	5.3	8.8	6.1
Final Extract pH (24 hr)	5.6	5.3	5.5	5.9
Acid Added (24hr)(ml)	406.4	2./	112.2	36.
Final Extract pH (28hr)	·	5.1	5.1	5.2
Acid Added (4hr)(ml)	_	2.	3/,	30.
Total Acid Added (ml)	406.4	4.1	143.2	66.
Total DI Water Added (ml)	1626.	2016.	1883.	1964.
Original Liquid Phase(ml)	0	0	0	0
Final Extract Volume (ml)	2032.4	2020./	2026.2	2030.
* * * * * * * * * * * * * * * * * * * *	· * * * * * *	* * * * *	* * * * * *	* * * * *
RCRA Metals				
Arsenic (PPM)	<0.01	<0.01	0.023	<0.01
Barium (PPM)	<0.5	< 0.5	1.0	1.1
Cadmium (PPM)	0.80	0.64	0,19	0./2
Chromium (PPM)	0.07	40.05	<0.05	<0.05
Lead (PPM)	0.45	<0.2	<0.2	<0.2
Mercury (PPM)	<0.001	<0.001	<0.001	<0.00
Selenium (PPM)	< 0.01	<0.01	<0.01	0,01
		1		

Note: Tables of pH adjustments with time have been extracted from the EIS sample work sheets and are reproduced in the section of this report containing laboratory bench sheets.

# ANALYTICAL REPORT SHEET SUPPLEMENTAL ANALYSIS - HAZARDOUS WASTES

This supplemental report sheet presents results of Corrosivity, Ingitability and Reactivity analysis if so required by the client. The following definitions apply to this analysis. The method of analysis is listed on the reverse side of this form.

- 1. Corrosivity A solid waste exhibits the characteristic of Corrosivity if it is aqueous and has a pH of less than (<) or equal to 2 or greater than (>) or equal to 12.5
- 2. Ignitability A solid waste exhibits the characteristic of ignitability if it is a liquid, other than an aqueous solution, containing less than 24% alcohol by volume, and has a Flash Point less than 140 °F (60 °C)
- 3. Reactivity Per procedures required by the Indiana State Board of Health, Reactivity is evaluated on the basis of Cyanide and Sulfide presence in the sample

Client Reactivity (ppm) \* Ignitability EIS Lab Corrosivity Sample Cyanide Sulfide (°F) (%C) (pH) Number Description Sample A - Comp. 6,4 9.4 > 158. 3.9 6516 Pretreatment was te Sample B-Paper 6.6. > 195. 3.9 5.1 6526 Filter, Production NO FLASH-EXTINGUISHES Sample C-Floor 3.1 5.8 9.1 6536 FLAME @ 140 Sweepings Sample D-Solid Paint waste 2.0 6.4 2.8 > 198 6546 200 760

\* Dry weight Basis

. 3

D. Corrective Action

### UNITED STATES ENVIRONMENTAL PROTECTION AGENCY REGION V

JUN 2 9 1987 Date:

Trip report of Visual Site Inspection (VSI) of VITCO (Formerly Vitreous Subject:

Steel Products Co.) IND 005081799

Herbert Levine, Geologist From:

Indiana Unit

Hak Cho, Chief To:

Indiana Unit

On June 29, 1987, I made the VSI for VITCO, Nappanee, Indiana. During the inspection I was accompanied by Mr. Brett Nordman representing VITCO. RCRA FACILITY ASSESSMENT FOR VITCO NAPPANEE, INDIANA INDO05081799

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U.S. EPA REGION V SOLID WASTE BRANCH TECHNICAL PROGRAMS SECTION JUNE 30, 1987

JUN 3 0 1987

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### Executive Summary

A RCRA Facility Assessment (RFA) was completed for VITCO, Nappanee, Indiana. The purpose of the RFA was to determine the need for corrective measures from any Solid Waste Management Unit (SWMU). The Hazardous and Solid Waste Amendments (HSWA) of 1984 require all RCRA permits issued after November 8, 1984, to contain provisions requiring corrective measures for all releases of hazardous waste or constituents from any SWMU at a treatment, storage, or disposal facility, regardless of when the waste was placed in such a SWMU. This facility decided to terminate their treatment operation and close its RCRA regulated SWMU before a final permit could be issued. The unit was clean-closed and received certification of closure from IDEM.

The RFA included a Preliminary Review (PR) and a Visual Site Inspection (VSI). A sampling Inspection (SI) was not required because the unit was clean-closed prior to the VSI. Since no other SWMUs exist that require corrective action a RCRA Facility Investigation (RFI) is not required.

### I. <u>Introduction</u>

The RCRA RFA has been completed for VITCO at Nappanee, Indiana. The purpose of the RFA was to determine the need for corrective action to address releases from any SWMU. The HSWA of 1984 require all RCRA permits issued after November 8, 1984, to contain provisions requiring corrective measures for a releases of hazardous wastes or constituents thereof.

This RFA includes both a PR and a VSI. The PR included a review of EPA files and a incomplete VSI begun but not completed by Mr. Gregor Weber of U.S. EPA, Technical Program Section, Indiana Unit.

### II. General Description

The VITCO facility is located in the town of Nappanne, Indiana. The facility is confined to two buildings; a general manufacturing building and a shop facility locate in another building on site. The general operations at the facility consists of metal forming and porcelain enamel application. The facility mainly applies porcelain paint. Prior to painting, metal parts are cleaned in a pickle-house by dipping in a acid bath. Over spill from this procedure enters a waste-water treatment unit. This unit generates a sludge that is handled as a hazardous waste and is transported off-site for disposal. The water from the waste-water treatment unit is discharged to the city sewer lines.

Observed Sc. veloase

### III. Solid Waste Management Units

### A. Waste-Pile Treatment Unit

This is the only RCRA regulated SWMU at the facility. This unit was clean-closed and has received certification of closure from IDEM.

### Discussion:

Visual Inspection revealed that no waste, container or residue remained.

### B. Waste-Water Treatment Unit

This unit treats spills that occur in the pickle degreasing house. A collection system collects all waste-water that is directed to the unit. Over-spray dust from the porcelain enamel painting operation is mixed with water and sent to this unit. The waste-water treatment unit generates a sludge that is seperated from the clarified water. The water is discharged to city sewer lines. This procedure is in compliance with the Water Division of IDEM. The sludge is collected and stored for subsequent off-site disposal.

### Discussion

Visual inspection revealed that no releases to the environment are occurring or have occurred from this unit.

### C. Container Storage

The facility maintains a roll-away bin for storage of sludge generated at the waste-water treatment unit. The bin is removed on a weekly basis for off-site disposal to a regulated facility.

### Discussion

The bin is well maintained, has a cover, and is stored on a cement pad. Visual inspection revealed that no releases had occurred.

### Conclusion

No further action is required for this facility. The only RCRA regulated unit has been clean-closed and received certification of closure by IDEM. During the VSI no additional SWMUs were observed. The facility will formally request withdrawal of their Part A application. Neither a SI or RFI is required for this facility.

### VISUAL SITE INSPECTION FOR WASTE MANAGEMENT UNITS

Facility Name EPA ID Number

Location Inspector

Title Facility Representative VITCO

IND 005081799 Nappanee, Indiana Herbert Levine

Geologist Brett Nordman

### 1. Facility General Description:

VITCO, located in Elkhart County, Indiana is a metal former and enamel porcelain applicator. Metal parts are cleaned in an acid bath and then painted with a enamel procelain paint. The facility applied for a RCRA permit by submittal of a Part A Permit Application. Only one RCRA regulated SWMU existed on site, a waste-pile treatment unit. This has been clean closed and the facility has received cerfitication of closure from IDEM.

### Site Description: II.

The facility is located on the out skirts of Nappanre. Adjacent to the facility is farm land. Residential houses are within 1/1/2 mile of the facility. No Part B application was submitted, there fore detailed geologic information is not available.

### III. Solid Waste Management Units:

- The facility had, in the past, operated a waste treatment unit. The facility closed this unit and has received a closure certification from the IDEM. During the VSI this unit was checked and noted to be closed.
- B. The facility operates a waste-water treatment unit. This unit receives waste-waters containing porcelain enamel dust and used acids from the pickle house. The waste water is treated with clarified water being discharged to city sewer lines and sludge collected for off-site disposal. The facility is in compliance with the Water Division of IDEM for their discharge. The facility has analysed the sludge and is asking the state to allow classification of the sludge as a special waste since it is not a characteristic waste.
- C. The facility maintains a roll-away container to receive sludge from the waste-water treatment unit. This bin is in good condition and is always kept covered. The bin is picked up each week so storage is not longer than 7 days.

### IV. General Observations:

There was no visual evidence of a release from any of the SWMUs. The only RCRA regulated unit (waste-pile treatment) has been certified closed.

### V. Conclusions:

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No corrective action is required for this facility. A RFI is not needed. This facility will formalley request withdrawal of their Part A application.

PRELIMINARYT REVIEW REPORT (PR) RCRA FACILITY ASSESSMENT (RFA)

	Vit	RCRA FACILITY ASSES	SSMENT (RFA)	*
	Facility Name (\	Jitreous Steel H 005081799 gor Weber	oducts (0.)	
2.	General Descript	cion of Facility and Processes		
	A. Description			
	B. Information	on Solid Waste Management Uni	ts (attach additional sheets as need	ed):
	B. Information	on Solid Waste Management Uni	ts (attach additional sheets as need Release (yes/no/unknown/suspec	
		Unit		
		u - 6		
	i. contra ii. treatm iii. waste	Unit		
	i. contra ii. treatm iii. waste iv.	unit unier storage ent bein, - closed		
	i. contra ii. treatm iii. waste iv.	Unit vier storage ent bin, - closed water treatment		
	i. contra ii. treatm iii. waste iv.	Unit vier storage ent bin, - closed water treatment	Release (yes/no/unknown/suspec	
	i. contra ii. treatm iii. waste iv. v. vi. vii.	Unit vier storage ent bin, - closed water treatment	Release (yes/no/unknown/suspec	
	i. contra ii. treatm iii. waste iv.	Unit vier storage ent bin, - closed water treatment	Release (yes/no/unknown/suspec	

Unit Type:	Regulatory Status:
Age:	
Capacity:	
Waste Type:	
Hazardous Constituents (attach sepa	rate sheet):
Mazaradas conscretenes (accaem sepa	7400 31100071
Unit Description:	
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Bulleten 9 Freezenskaan Noododo	
Additional Information Needed:	
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	,

Additional Information Needed:		vescription	(groundwater,	surrace water,	, etc.):	
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OFFICIAL PHOTOGRAPH

U.S. ENVIRONMENTAL PROTECTION AGENCY

PROJECT/CASE NO. TND OOS O 8 | 799

SUBJECT: 1-00

LOCATION: 900 E. Wa bash Ave

LOCATION: 900 E. Wa bash Ave

DATE: 2-1287 TIME:

WEATHER SUN (HAZE) ICLOUDY (RAIN) (SNOW)

WITNESS:

CAMERA: VIKON FG SO NOW LONG

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10 JN: 900 E. Wahash
CHAPPANER COUNTY Elkhast STATE IND
DATE 2-12-87 TIME
DATE 2-12-87 TIME
WEATHER: (SUM PHAZE) (CLOUDY) (RAIN) (SHOW)
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OFFICIAL PHOTOGRAPH

U.S. ENVIRONMENTAL PROTECTION AGENCY

PROJECT/CASE NO: VITCO

SUBJECT: IND OCISOBIZ 99

LOCATION: 900 E Wa Dash

NCITY! Dan eccounty! (Chart STATE IND

DATE I 12 87 TIME

WEATHER (SUM CHARE) (COUNTY) GRAIN (SNOW)

PHO UPHER (SIE)

WITI.

CAMERA: NIKON FG SO MAN ICM S

FILM TYPE 12 6 ASA ZOTO 1/ F

NEGATIVE LOCATION: FILE IF

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OFFICIAL PHOTOGRAPH U.S. ENVIRONMENTAL PROTECTION AGENCY PROJECT/CASE NO; IND 005 08 1799
SUBJECT OF TEO SUBJECT VITO Nappanee COUNTY & Ikhart STATE IND

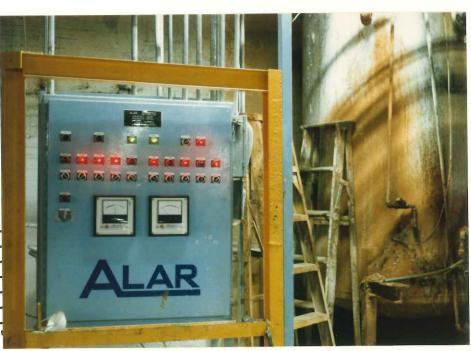
DATE 2-12-87 TIME

DATE 2-12-87 TIME WEATHER: (SUM) PHAZE (CLOUDY) PRAING SHOWN PHOTOGRAPHER (SIL) CAMERA: NIKON FR SOMM FILM TYPE 126 ASIZOO T:1/ NEGATIVE LOCATION: FOA of warter makes 800 391-399 PHOTO #: \_\_



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U.S. ENVIRONMENTAL PROTECTION AGENCY SUBJECT/CASE NO: IND 005081799
SUBJECT. VII CO
LOCATION 900 E Wabash Nappanee county & | chaf STATE IND

DATE 2-12-87 TIME WEATHER ISUNG PHAZE ICLOUON PRAINS, ISNOWN, PHOTOGRAPHER (SIE) CAMERA: NI WON FG SU NAME \_1312UD 11/\_ FILM TYPE TYL MEGATIVE LOCATIONS EOA) PROCESSED 8Y: Collection of Pit for

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OFFICIAL PHOTOGRAPH U.S. ENVIRONMENTAL PROTECTION AGENCY SUBJECT VI TED LOCATION: 900 E Nappanee county Elehart STATE IND

DATE 2-12-87 TIME WEATHER: (SUM) CHAZE CLOUDY, FRAIN SNOW) CAMERA NIKON F.G. 50 mm FILM TYPE 17.6 ASUZO OT:1/\_ PROCESSED BY: . Caustic-cleener holding PHOTO #: -

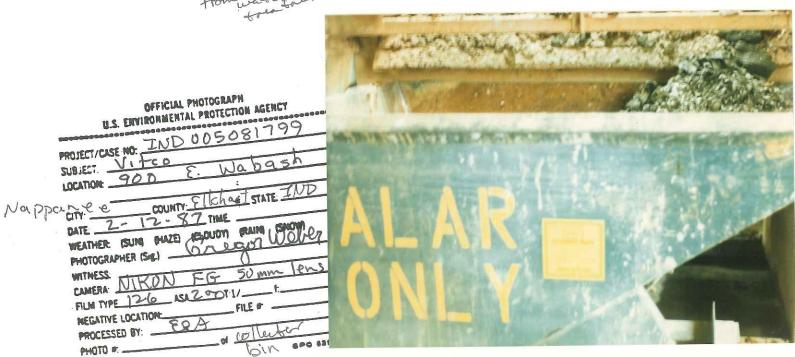


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SUBJECT: VI + CO
LOCATION: 900 E Wahash DATE 2-12 87 TIME Nappartee WEATHER ISUN (HAZE) GLOUDY (RAIN) SNOW PHOTOGRAPHER (SIE) CAMERA: NIKON FG 50 mm lens FILM TYPE: 12 G ASAZO OT:1/. NEGATIVE LOCATION\_ EDA PROCESSED BY: \_\_\_ Depo 138 - 130 PHOTO . fromwadownt

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W.S. ENVIRONMENTAL PROTECTION AGENCY

PROJECT/CASE NO; INDOOS 081799

PROJECT/CASE NO; INDOOS 081799

SUBJECT 900 E. Wabash

LOCATION: 900 E. Wabash

CITY! 2-1287 TIME

PROTOGRAPHER (SEL) - WATER CLOUDY PRAIM SNOWY

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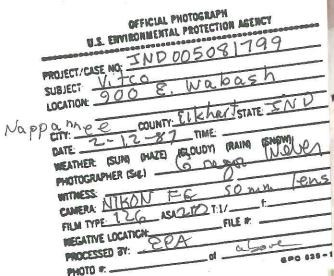
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U.S. ENVIRONMENTAL PROTECTION AGENCY
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DATE: 2 - 12 - 8 7 TIME: WEATHER: (SUN) (HAZE) (GLOUDY) (RAIN) (SNOW) PHOTOGRAPHER (SIE)	
WITNESS:  CAMERA: VIKO V FG 50 mm   Ph    FILM TYPE 12 G ASAZOO T:1/ f:  NEGATIVE LOCATION: FILE #:  PROCESSED BY: EPA	S
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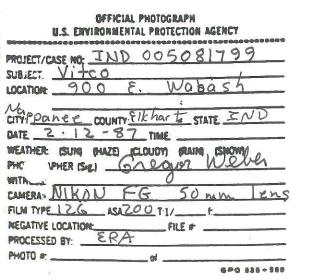
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PROJECT/CASE NO: IND 005081799
SUBJECT. VITCO
Na pounce county Elkhart STATE IND
DATE 2-12-87 TIME WEATHER: (SUNG PHAZE) (CLOUDY) (RAING (SHOW))
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U.S. ENVIRONMENTAL PROTECTION AGENCY
PROJECT/CASE NO: IND 005081799
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DATE 2-12-87 TIME
WEATHER: (SUM) (HAZE) (CLOUDY) (RAIM) (SHOW) PHOTOGRAPHER (Sig.) — (SHOW)
CAMERA: NIKON FG 50 mm lens
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CASE NO: VITCO
SUBJECT TND OBJUSTASA
LOCATION: SOO
Vappanneeounty 11 Khat STATE IND
DATE 7 - 17 - 8 / HINE BAING SNOW
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OFFICIAL PHOTOGRAPH U.S. ENVIRONMENTAL PROTECTION AGENCY PROJECT/CASE NO: TND
SUBJECT VITCO LOCATION: 900 Nagganneecounty Fikhai State IA DATE - 2 - 12 - 8 WEATHER ISUNG CHAZE ACCOUDY (RAIN) PHOTOGRAPHER (SIE) -CAMERA: MKON WITNESS: Summ FILM TYPE 1 7 Co ASA 7 OOT:1/ NEGATIVE LOCATION PROCESSED BY: \_ or 011 PHOTO #: -



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Name of Preparer: Gregor Weber.
Date: 02-11-87

Model	Faci	lity	Management	Plan
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Viteo

- . 1. Facility Name: Vitreous Steel Products Co.)
  - 2. Facility I.D. Number: <u>IND 005 081799</u>
  - 3. Owner and/or Operator: Vitreous Steel Prod. Co.
  - 4. Facility Location: 900 East Wabash Ave.

    Street Address

Nappanee Elkhart Indiana 46550 Hoity County State Zip Code

- 5. Facility Telephone (if available): (219) 773 318/
- 6. Interim Status and/or Permitted Hazardous Waste Units and Capacities of Each Unit:

Type of Units Size or Capacity Active or	Closed
Storage in Tanks or Containers	
Incinerator	
Landfill	
Surface Impoundment	
Waste Pile	•
Land Treatment	8.
Injection Wells	
TO4 Others (Specify) 450 liters/day Active (Treatment Bin) Close of	
7. Permit Application Status:(HVDMS action item number)	

8.	Identification of Hazardous Waste Generated, Treated, Stored or Disposed at the Facility: ( may attach Part A or permit list or reference those documents if listing of wastes is
<b>8</b>	exceptionally long - in that case, to complete this question list wastes of greatest interest and/or quantity and note that additional wastes are managed)

Type of Waste Quantity Generated, Treated, Stored or Disposed (note appropriate categories)

Paint waste EP Toxic (D005, D006) Generated &

186 Ton/Year

9. Review of Response to Solid Waste Management Questionaire indicates: (check one)
waste Management Questionaire indicates: (check one)
identified RCRA units)
No Solid Waste Management Units exist (other than previously identified RCRA units)
It is unclear from review of questionaire whether or not any solid Waste Management Units exist
Respondent indicates that does not know if any Solid Waste
10. If the response to question 9 is that Solid Waste Management Units exist, than check one of the following:
Releases of hazardous waste or constituents have occurred or are thought to have occurred
Releases of hazardous waste or constituents have not occurred
Releases of hazardous waste or constituents have occurred or are thought to have occurred but have been adequately remedied
It is not known whether a release of hazardous waste or constituents has occurred

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Yes - indicate List or update	•
No	
Yes - ERRIS list	
n of the Recommendation portion of the Facility Managem ndix must be completed.	21
Regional Approach to the Facility: Check one	
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ion Order (may include compliance schedule)	
rative Enforcement	
al Enforcement	
RCLA for Federally Financed or Enforcement Activity	
tiated Action	
anation of selection:	
tion alternative is selected:	
on - anticipated inspection date	
State or Federal inspection	
Assessment - anticipated completion date	
ipated date of initiation	
State/Federal	
double name identific name (ide)	
Private Party identify party(ies)	
	Yes - indicate List or update  NO  Yes - ERRIS list  of the Recommendation portion of the Facility Management and the completed.  Regional Approach to the Facility: Check one agation to Evaluate Facility  noce Schedule  Ion Order (may include compliance schedule)  reative Enforcement  ALLA for Federally Financed or Enforcement Activity  triated Action  anation of selection:  State or Federal inspection  Assessment - anticipated completion date  ipated date of initiation  State/Federal

b)	If Permit Alternative is Selected: Projected Schedule
	Date of Part B Submission:
	Date of Completeness Check:
•	Date for Additional Submissions (if required):
•	Date of Completion of Technical Review:
٠	Completion of Draft Permit/Permit Denial:
	Public Notice for Permit Decision:
	Date of Hearing (if appropriate):
	Date for Final Permit or Denial Issuance:
	Description of any corrective action provisions to be included in permit -
c)	If Corrective Action Order Alternative is Selected:
	Estimated Date for Order Issuance:
	Description of Provisions of the Order to be Completed by Facility:
	Description of Compliance Schedule to be Contained in Order:
	/
d)	If Other Administrative Enforcement Action is Selected:
	Projected Date for Issuance of the Order:
	Description of Provisions or Goals of the Order:

If Judicial Enforcement Alternative Selected:
Date of Referral to Office of Regional Counsel:
) If Referral to CERCLA for Action Selected:
Date of Referral to CERCLA Sections:
) If Voluntary/Negotiated Action Alternative if Selected:
Date of Initial Contact with Facility:
Description of Goals of Contact or Discussions with Facility:
Date for Termination of Discussions if Not Successful:
Date of Finalization of Settlement if Negotiation Success
h) If State Action Alternative is Selected:
Date for Referral to State:
Name of State Contact:
Phone:

### APPENDIX

The questions constituting this Appendix to the Facility Management Plan must be filled out prior to completion of recommendation elements of the Plan. The purpose of this appendix is to provide a summary documentation of the State and/or U.S.EPA review of available information on the subject facility. The intent is that a comprehensive file review will be conducted as the basis for selection of the recommended approach to a given facility. If the Appendix is information in State personnel questions referring to available data reference files. Where questions refer to "all" available data or information and such material is voluminous, the response should indicate that files are voluminous, and then reference most telling information, for example groundwater contaminants found and information most directly supporting recommended approach to facility should be described. If no information is available in facility files, the response should so indicate. It is also anticipated that this Appendix may be updated periodically as more information becomes available.

1. Description of All Available Monitoring Data for Facility:

Type of Data Date Author Summary of Results or Conclusions

2. Description of Enforcement Status:

Type of Action Date Local, State or Federal Result or Status

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3. Description of Any Compl	laints from Publi	C:	
Source of Complaint	Date Recipient	Subject and Response	
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4. Description of All Inspe	ection Reports fo	r Facility:	
Date of Inspection	Inspector (Local Fe	,State, Conclusions or deral)	Comments
		·	
disposal practices no	ot currently regul n wells. ponds or	the inspector note any evidence lated under RCRA such as pil surface impoundments that mandfills?	es of waste
Yes - gi	ive date if inspe	ction and describe observati	ion
***************************************			<u> </u>
-		•	•
No		·Don¹t	know

No

6. Do inspection reports indicate observations of discolored soils or dead vegeta- tion that might be caused by a spill, discharge or disposal of hazardous wastes or constituents?
Yes - indicate date of report and describe observations
No No
Don*t know
7. Do inspection reports indicate the presence of any tanks at the facility which are located below grade and could possibly leak without being noticed by visual observation?
Yes - date of inspection and describe information in report
No
Don't know
8. Does a groundwater monitoring system exist at the facility?
9. If answer to question 8 is yes, is the groundwater system capable of monitoring both regulated RCRA units and other Solid Waste Management Units?
Explain -
10. Is the groundwater monitoring system in compliance with applicable RCRA groundwater monitoring standards?
If no, explain deficiency

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Type of Information Author Date Summary of Conclus  12. Did the facility submit a 103(c) notification pursuant to CERCIA?  Yes Date of Notification  No  13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been confort this facility?  Yes  No	11.		all informati lable.	ion on fac	ility subs	urface ge	eology or hydrogeology
12. Did the facility submit a 103(c) notification pursuant to CERCIA?  Yes Date of Notification  No  13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been confor this facility?  Yes  No		Type of	Information	Author	Date		Summary of Conclus
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Yes Date of Notification  No  13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been confor this facility?  Yes  No			,				
Yes Date of Notification  No  13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been confor this facility?  Yes  No							
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13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCLA Preliminary Assessment/Site Investigation (PA/SI) been confor this facility?  Yes  No	12.	Did the	e facility sub	mit a 103(	c) notific	ation pu	rsuant to CERCLA?
13. If answer to 12 is yes, briefly summarize content of that notification (waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been confor this facility?  Yes  No			Yes	Date of	Notificat	ion	
(waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been corfor this facility?  Yes  No			No				
(waste management units identified, type of waste concerned)  14. Has a CERCIA Preliminary Assessment/Site Investigation (PA/SI) been corfor this facility?  Yes  No							
(waste management units identified, type of waste concerned)  14. Has a CERCLA Preliminary Assessment/Site Investigation (PA/SI) been corfor this facility?  Yes  No	13.	If answ	wer to 12 is y	es, brief]	y summariz	e conter	nt of that notification
for this facility?  Yes  No		()	waste manageme	ent units i	dentified,	, type of	waste concerned)
for this facility?  Yes  No							,_
for this facility?  Yes  No							
for this facility?  Yes  No							
for this facility?  Yes  No				-			
for this facility?  Yes  No						_	
Yes No	14. F	las a CE or this	RCLA Prelimina facility?	ary Assess	ment/Site 1	Investiga	ation (PA/SI) been com
No .		•					
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	cusing on types of environmental contamination found, wastes and source contamination.
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l6. If a	available, having reviewed the CERCLA notification, RCRA Part A and RC B, it appears that: (CERCLA unit refers to unit or area of concern
16. If a	CERCLA response activity)
16. If a	CERCLA response activity)  RCRA and CERCLA units are same at this facility
16. If a	RCRA and CERCLA units are same at this facility  RCRA and CERCLA units are clearly different units
l6. If a	CERCLA response activity)  RCRA and CERCLA units are same at this facility  RCRA and CERCLA units are clearly different units  There is an overlap between the
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16. If a	CERCLA response activity)  RCRA and CERCLA units are same at this facility  RCRA and CERCLA units are clearly different units
16. If a	CERCIA response activity)  RCRA and CERCIA units are same at this facility  RCRA and CERCIA units are clearly different units  There is an overlap between the

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18. Identification of Reports or Documentation Concerning Each Release Described in Item 17.

Title/Type of Report Date Author Recipients Contents

19. Highlight any information gaps in the file - describe any plans to obtain additional needed information.

20. Summary of major environmental problems noted, desired solution and possible approaches.

Problem Solution Approach Pros and Cons





### VITREOUS STEEL PRODUCTS COMPANY

900 EAST WABASH AVENUE · NAPPANEE, INDIANA 46550

PHONE (219) 773-3181 · TELEX 25-8351

FEB 2 4 1986

U.S. EPA, REGION V

February 20, 1986

RCRA Activities Region V P.O. Box A3587 Chicago, IL 60690

Attention: Atkjg

Gentlemen:

Attached is the certification regarding potential release from solid waste management units.

Yours truly,

VITREOUS STEEL PRODUCTS

John F. Kendall

JFK/nsm

Attachment

# CERTIFICATION REGARDING POTENTIAL RELEASES FROM SOLID WASTE MANAGEMENT UNITS

FACILITY NAME:	VITREOUS STEEL PRODUCTS COMPANY
EPA I.D. NUMBER:	IND 005081799
LOCATION CITY:	Nappanee
STATE:	Indiana 46550
التحيد فعالهممماهم	of the following solid waste management units (existing or facility? NOTE - <u>DO NOT INCLUDE HAZARDOUS WASTE UNITS</u> N IN YOUR PART A APPLICATION
Storage Tan Container S Injection W Wastewater Transfer St Waste Recycl Waste Treat Other  If there are provide a desc	k (Above Ground)  k (Underground)  torage Area  ells  Treatment Units  ations  ling Operations ment, Detoxification  Yes" answers to any of the items in Number 1 above, please or iption of the wastes that were stored, treated or disposed
would be cons RCRA. Also in disposed of a	idered as hazardous wastes or hazardous constituents under not lude any available data on quantities or volume of wastes and the dates of disposal. Please also provide a description and include capacity, dimensions and location at facility. e plan if available.
* Container sto	rage area - Less than 90 day storage
* Waste water t	reatment system - Installed 25 Nov. 85 to comply with
regulations f	or point source porcelain enamelers. This system is
exempt from tr	eat store and dispose requirements.
	<b>\$</b> **
NOTE: Hazaro	dous wastes are those identified in 40 CFR 261. Hazardous

to the environment occurring.						
Please provide th	ne followin	g informat	ion			٠
a. Date of releads b. Type of waste c. Quantity or of d. Describe nation or tank, etc	e released volume of w ure of rele	aste relea ase (i.e.,	sed spill, ove	erflow, rup	tured pi	pe
No known releas	ses					- Contract of the Contract of
	1 .			. •		
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which would desc that exists as a hazardous wastes	ribe the na result of or constit	sture and e such relea	extent or e ises. Plea	nvironmenta se focus of	n concent	ratio
which would desc	ribe the na result of or constit	sture and e such relea	extent or e ises. Plea	nvironmenta se focus of	n concent	ratio
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please provide ( which would desc that exists as a hazardous wastes  Not applicable	ribe the na result of or constit	sture and e such relea	extent or e ises. Plea	nvironmenta se focus of	n concent	ratio
which would desc that exists as a hazardous wastes	ribe the na result of or constit	sture and e such relea	extent or e ises. Plea	nvironmenta se focus of	n concent	ratio
Not applicable  I certify under prepared under resigned to assist information who manage the information true, accurate, ties for submit and imprisonmen	penalty of my directioure that quested system, or the submiting false to for knowing the submiting false to the su	law that n or super alified pe Based on those pers ttal is, t	this docume vision in a rsbnnel pro my inquir ons direct o the best aware that n includi	ent and all accordance operly gath y of the pely responsi of my know there are no the poss	attachme with a sy er and er rson or p ble for ledge an signification	ents wystem valuat person deliant person fir
I certify under reprepared under resigned to assist information who manage the information true, accurate,	penalty of my directioure that quested system, or the submiting false to for knowing the submiting false to the su	law that n or super alified pe Based on those pers ttal is, t	this docume vision in a rsbnnel pro my inquir ons direct o the best aware that n includi	ent and all accordance operly gath y of the pely responsi of my know there are no the poss	attachme with a sy er and er rson or p ble for ledge an signification	ents wystem valuat person deliant person fir

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## Site Investigation Report

1)	During the inspection of this facility did the inspector note any evidence of past waste disposal practices not currently regulated under RCRA such as piles of waste or rubbish, ponds or surface impoundments that might contain waste, active or inactive landfills?
	a) Yes, Explain
	b) No
	c) Cannot Respond to this Ouestion
2)	Was there any evidence of discolored soils or dead vegetation that might be caused by a spill, discharge or disposal of hazardous wastes or constituents?
	a) Yes, Explain
	b) No
	c) Cannot Respond to this Question
3)	(solid or hazardous) which are located below grade and could possibly leak without being noticed by visual observation? a) Yesb) No
	c) Cannot Respond to this Question
4)	Based on an inspection or inspections that have been done at this facility there is no reason to question or doubt the information which the applicant has submitted on the questionnaire regarding Solid Waste Management Units and the possibility of prior or continuing releases of hazardous wastes or constituents.
	a) I concur with this statement
	b) I do not concur with this statement for the following reasons:

5)	If 4(b) was checked, Describe what additional information or testing is needed to determine if prior or continuing releases of hazardous wastes or constituent have occurred. Specify which units are of concern and what types of releases are suspected (i.e., releases to groundwater, surface water, air, soils, etc).
6)	An on site inspection to discuss and evaluate the possibility of prior or continuing releases from Solid Waste Management Units is recommended
	a) Yes
	<u>×</u> b) No
7)	Was site sampling for confirmation of suspected releases conducted?
	a) Yes
	b) No
8)	If yes to 7, detail the following:
	<ol> <li>Sampling plan - include locating parameters to be tested. rationale for each parameter, logistics, dates, personnel etc.</li> </ol>
•	2) Analytical results - QA, QC. Result summary conclusion.
9)	A Remedial Investigation (R.I.) is needed to evaluate the nature and extent of prior releases of hazardous wastes or constituents from Solid Waste Management Units.
	a) Yes
	b) No

10.	If the answer to No. 7 above is "Yes", the priority and manner for requiring the applicant to conduct the Remedial Investigation (R.I.) is as follows:
	a) Require R.I. in compliance schedule that is part of RCRA permit.
	b) Issue Compliance Order requiring R.I. to be done.
11.	Did the SI address all items that the PA "Assessment of Unit" forms indicated the SI should address?
	y y y
12.	Based on my review of this S.I. report, it is hereby:
	not approved  Signature Act Le Date: 6-29-87  (EPA Staff)

Facility Vitreaus Steel I.D. No. IND 045081799

Page No. A.

	STE MANAGEMENT UNITS AND MAJOR SPILLS						
Unit or Spill		OCATI	0 N	0 F	NFOR		1 0 N
	Permit Applic	SWMU Questnr	NPDES Files	Enfrcmt Files	CERCLA Files	State Files	Other
1. Container storage	X						RFA
2. treatment bin - closed	× _						KFA
3. Waste-water treatment	×					· · · · · · · · · · · · · · · · · · ·	RFA File
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SUBJECT B. GENERAL FACILITY DESCRIPTION

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Catego	ry Descri	ntion	0.4			
1		ON/SETTING/LAND USE/SIZE	Category	Description		
2	TNOUST	RY TYPE	<u> </u>	OPERATIONAL HISTORY		
3	PPODIIC	TS PRODUCED	REGULATORY STATUS OTHER			
	DAU MA	TERIALS				
, ,	<u> </u>	TERTALS	8			
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	5	This facilit	<u> </u>	1	Part A	
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7				plière	Entorcement	
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SUBJECT C. ENVIRONMENTAL SETTING, HYDROLOGICAL AND GEOLOGICAL CHARACTERISTICS	SUBJECT C.	ENVIRONMENTAL S	SETTING,	HYDROLOGICAL	AND GEOL	DGICAL	CHARACTERISTICS
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1	UBJECT C.	ENVIRONMENTAL SETTING, HYD	ROLOGICAL	AND GEOLOGICAL	L CHARACTERISTICS
Catego	ry Descrip	tion	Category	Description	
1	GEOLOGY		7	SOIL DATA	
2				ATMOSPHERIC	DATA
3	WELL LOCA	ATION/CONSTRUCTION OFF-SITE	9	PHOTOGRAPHS/	MAPS
4	GROUNDWA	TER MONITORING DATA	10	OTHER	
5	SURFACE I	71,000			
6	FLOODPLA	IN DATA	12		
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ITEM	CATEGORY	DESCRIPTION/DATES/	COMMENTS		LOCATION OF INFORMATION
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		rule of face	lity		
	4	none	7		
	_5	none			
	6	no			
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	8	none			
··	9	Plotos of fa	ci li ly	wer	RFA file
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SUBJECT D. WASTE CHARACTERIZAT	TON
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Category	Descr	iption		Category	Description			
1			RCRA IDENTIFIED	4	WASTES - STA	TE IDENTIFIED		
<u>la</u>			/Constituents	onstituents 4a Descripti				
<u>lb</u>	Amo	unts pla	ced/spilled - Date	<b>4</b> b		laced/released		
2			NFORCEMENT IDENTIFIED	5	INSPECTIONS			
Ža	Des	cription	/Constituents	6	STATE PERMITS			
<b>2</b> b	Алто	unts pla	ced/released	ed/released 7 WASTES - F				
3			CLA IDENTIFIED 8 WASTES - OTHER IDENTIFIED					
3a			/Constituents	9	OTHER			
<b>3</b> b	Amo	unts pla	ced/released					
++++++	++++++	++++++	+++++++++++++++++++	+++++++	+++++++++	<del></del>		
ŀ	ATEGORY	SPILL	DESCRIPTION/DATE			LOCATION OF INFORMATION		
		#2	Treat ment	Bin		Part A		
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SUBJECT E. UNIT/SPILL SPECIFIC INFORMATION	
UNIT/SPILL: Treatment Bin	

Catego	ry Descrip	tion Category Descr	iption
1	ENGINEER	110 DECEMBER 1 DESCRIPTION OF THE PROPERTY OF	INSPECTIONS
2	[WASTES	7 DUDIT	C SUPPLIED INFORMATION
3	OPERATIO	NAL STATUS/DATES OF USAGE 8 LOCAT	ION
4	PERMITS	Q ATUED	
5	TADEQUACY	TO PREVENT RELEASES	
*****	*********	++++++++++++++++++++++++++++++++++++++	<del>+++++++++++++++++++++++++++++++++++++</del>
ITEM	CATEGORY		LOCATION OF
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		1 3	RFA:
		Ouring G.W. 2-11-87 V	St file
	2	EPTOLIC (BOOS DOOG)	1 1 af
		Paint Waste	- Comment
	3	Closed as of 4-10-85	Path
	4	Submitted Pat A	PartA
	_5	closed as first S.	100411
	(e	RFA 119 1 10110	21 0 50 0
		no of care	2 kg G. W. RFA f
	8		
		inde conton Have &'Stock	Kroon Part H
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SUBJECT E. UNIT/SPILL SPECIFIC INFORMATION	
UNIT/SPILL: Worker Vreature +	
Category Description	
2 WASTES DESCRIPTION/PHOTOGRAPHS 6 RCRA INSPECT	IONS
3 OPERATIONAL STATUS/DATES OF USAGE 8 LOCATION	IED INFORMATION
5 ADEQUACY TO POEVENE 9 OTHER	
**************************************	**** <b>*</b> ***
DESCRIPTION/DATES/COMMENTS	LOCATION OF INFORMATION
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SUBJECT E. UNIT/SPILL SPECIFI	C INFORMATION	
UNIT/SPILL: Condainer Store	al	
Category Description	7	
1 ENGINEERING DESCRIPTION/PHOTOGRAPHS 6	y Description RCRA INSPECT	LONG
3 OPERATIONAL STATUS/DATES OF USAGE 7	PUBLIC SUPPL	IED INFORMATION
5 IADEQUACY TO ODE VENE	OTHER	
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The state of the s	SUBJECT F.	KNOWN &	SUSPECTED	RELEASES &	<b>EXPOSURE</b>	POTENTIAL	
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Cate-		Cate-	
gory	Description	gory	Description
1	GROUNDWATER	5	CONTAMINATED SOLIDS DISPERSION
la	Actual releases	5a	Actual releases
lb	Potential releases	<b>5</b> b	Potential releases
lc	Pathways	5c	Pathways
ld	Potential/actual Exposure (human)	5d	Potential/actual Exposure (human
le	Activities affected by a release	5e	Activities affected by a release
2	SURFACE WATER	6	TRANSPORTATION RELATED
2a	Actual releases	6a	Actual releases
<b>2</b> b	Potential releases	6b	Potontial palessas
2c	Pathways	6c	Potential releases
2d	Potential/actual Exposure (human)	6d	Pathways
2e	Activities affected by a release		Potential/actual Exposure (human)
3	AIR	<u>бе</u>	Activities affected by a release
3a	Actual releases		FOOD CHAIN CROPS
3b	Potontial calculation	7a	Actual releases
3c	Potential releases	7ь	Potential releases
3d	Pathways	7c	Pathways
	Potential/actual Exposure (human)	7d	Potential/actual Exposure (human)
3e	Activities affected by a release	7e	Activities affected by a release
4	SUBSURFACE GAS	8	PUBLIC COMPLAINTS/CONCERNS
4a	Actual releases	9	OTHER
<b>4</b> b	Potential releases		
4c	Pathways		
<b>4</b> d	Potential/actual Exposure (human)		
4e	Activities affected by a release		
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ITEM	CATEGORY	UNIT/ SPILL	DESCRIPTION/DATES/COMMENTS	LOCATION OF INFORMATION
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